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Patched-2 Antibodies

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RELATED APPLICATIONS

This application is a continuation of USSN 09/293,505 filed 15 April 1999, now allowed U.S. Pat. No. 6,348,575, which claims priority under 35 U.S.C. § 119(e) to provisional application no. 60/081,884, filed 15 April 1998, all of which the entire disclosure is hereby incorporated by reference.

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FIELD OF THE INVENTION

The present invention relates generally to signaling molecules, specifically to signaling and mediator molecules in the *hedgehog (Hh)* cascade which are involved in cell proliferation and differentiation.

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BACKGROUND OF THE INVENTION

Development of multicellular organisms depends, at least in part, on mechanisms which specify, direct or maintain positional information to pattern cells, tissues, or organs. Various secreted signaling molecules, such as members of the transforming growth factor-beta (TGF- β), Wnt, fibroblast growth factors and hedgehog families have been associated with patterning activity of different cells and structures in *Drosophila* as well as in vertebrates. Perrimon, *Cell*: 80: 517-520 (1995).

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Segment polarity genes were first discovered in *Drosophila*, which when mutated caused a change in the pattern of structures of the body segments. These changes affected the pattern along the head to tail axis. *Hedgehog (Hh)* was first identified as a segment-polarity gene by a genetic screen in *Drosophila melanogaster*, Nusslein-Volhard *et al.*, *Roux. Arch. Dev. Biol.* 193: 267-282 (1984), that plays a wide variety of developmental functions. Perrimon, *supra*. Although only one *Drosophila Hh* gene has been identified, three mammalian *Hh* homologues have been isolated: Sonic *Hh* (*Shh*), Desert *Hh* (*Dhh*) and Indian *Hh* (*Ihh*), Echelard *et al.*, *Cell* 75: 1417-30 (1993); Riddle *et al.*, *Cell* 75: 1401-16 (1993). *Shh* is expressed at high level in the notochord and floor plate of developing vertebrate embryos, and acts to establish cell fate in the developing limb, somites and neural tube. *In vitro* explant assays as well as ectopic expression of *Shh* in transgenic animals show that *Shh* plays a key role in neural tube patterning, Echelard *et al.* (1993), *supra*; Ericson *et al.*, *Cell* 81: 747-56 (1995); Marti *et al.*, *Nature* 375: 322-5 (1995); Roelink *et al.* (1995), *supra*; Hynes *et al.*, *Neuron* 19: 15-26 (1997). *Hh* also plays a role in the development of limbs (Krauss *et al.*, *Cell* 75: 1431-44 (1993); Laufer *et al.*, *Cell* 79, 993-1003 (1994)), somites (Fan and Tessier-Lavigne, *Cell* 79, 1175-86 (1994); Johnson *et al.*, *Cell* 79: 1165-73 (1994)), lungs (Bellusci *et al.*, *Develop.* 124: 53-63 (1997) and skin (Oro *et al.*, *Science* 276: 817-21 (1997). Likewise, *Ihh* and *Dhh* are involved in bone, gut and germinal cell development, Apelqvist *et al.*, *Curr. Biol.* 7: 801-4 (1997); Bellusci *et al.*, *Dev. Suppl.* 124: 53-63 (1997); Bitgood *et al.*, *Curr. Biol.* 6: 298-304 (1996);

Roberts *et al.*, *Development* 121: 3163-74 (1995). Specifically, *Ihh* has been implicated in chondrocyte development [Vortkamp, A. *et al.*, *Science* 273: 613-22 (1996)] while *Dhh* plays a key role in testis development. Bitgood *et al.*, *supra*. With the exception of the gut, in which both *Ihh* and *Shh* are expressed, the expression patterns of the hedgehog family members do not overlap. Bitgood *et al.*, *supra*.

5 At the cell surface, *Hh* function appears to be mediated by a multicomponent receptor complex involving *Patched* (e.g., *Ptch*) and *Smoothed* (e.g., *Smo*), two multi-transmembrane proteins initially identified as segment polarity genes in *Drosophila* and later characterized in vertebrates. Nakano *et al.*, *Nature* 341: 508-513 (1989); Goodrich *et al.*, *Genes Dev.* 10: 301-312 (1996); Marigo *et al.*, *Develop.* 122: 1225-1233 (1996); van den Heuvel, M. & Ingham, P.W., *Nature* 382: 547-551 (1996); Alcedo, J. *et al.*, *Cell* 86: 221-232 (1996);
10 Stone, D.M. *et al.*, *Nature* 384: 129-34 (1996). Upon binding of *Hh* to *Patched*, the normal inhibitory effect of *Patched* on *Smo* is relieved, allowing *Smo* to transduce the *Hh* signal across the plasma membrane. It remains to be established if the *Patched/Smo* receptor complex mediates the action of all 3 mammalian hedgehogs or if specific components exist. Interestingly, a second murine *Patched* gene, *Patched-2* was recently isolated [Motoyama, J. *et al.*, *Nature Genetics* 18: 104-106 (1998)], but its function as a *Hh* receptor has not been
15 established. In order to characterize *Patched-2* and compare it to *Patched* with respect to the biological function of the various *Hh* family members, Applicants have isolated the human *Patched-2* gene. Biochemical analysis of *Patched* and *Patched-2* show that both bind to all members of the *Hh* family with similar affinity and that both molecules can form a complex with *Smo*. However, the expression patterns of *Patched-2* and *Patched* do not overlap. While *Patched* is expressed throughout the mouse embryo, *Patched-2* is found mainly in spermatocytes
20 which require Desert Hedgehog (*Dhh*) for proper development suggesting that *Patched-2* mediates *Dhh*'s activity in the testis. Chromosomal localization of *Patched-2* places it on chromosome 1p33-34, a region deleted in some germ cell tumors, raising the possibility that *Patched-2* may be a tumor suppressor in *Dhh* target cells.

SUMMARY OF THE INVENTION

25 In one embodiment, the invention provides an isolated nucleic acid molecule having at least about 80% sequence identity to (a) a DNA molecule encoding a *Patched-2* polypeptide comprising the sequence of amino acids 1 to 1203 of Fig. 1 (SEQ ID NO:2), or (b) the complement of the DNA molecule of (a); and encoding a polypeptide having *Patched-2* biological activity. The sequence identity preferably is > 91%, more preferably about 92%, most preferably about 95%. In one aspect, the isolated nucleic acid has at least > 91%, preferably at least about 92%, and even more preferably at least about 95% sequence identity with a polypeptide having amino acid residues 1 to about 1203 of Fig. 1 (SEQ ID NO:2). In a further aspect, the isolated nucleic acid molecule comprises DNA encoding a human *Patched-2* polypeptide having amino acid residues 1 to about 1203 of Fig. 1. In yet another aspect, the invention provides for an isolated nucleic acid comprising DNA having at least a 95% sequence identity to (a) a DNA molecule encoding the same mature polypeptide encoded by the cDNA in ATCC Deposit No. 209778 (designation: pRK7.hptc2.Flag-1405), alternatively the coding sequence of clone pRK7.hptc2.Flag-1405, deposited under accession number ATCC 209778. In a still further aspect, the
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invention provides for a nucleic acid comprising human *patched-2* encoding sequence of the cDNA in ATCC Deposit No. 209778 (designation: pRK7.hptc2.Flag-1405) or a sequence which hybridizes thereto under stringent conditions.

5 In another embodiment, the invention provides a vector comprising DNA encoding a human *patched-2* polypeptide. A host cell comprising such a vector is also provided. By way of example, the host cells may be mammalian cells, (e.g. CHO cells), prokaryotic cells (e.g., *E. coli*) or yeast cells (e.g., *Saccharomyces cerevisiae*). A process for producing *patched-2* polypeptides is further provided and comprises culturing host cells under conditions suitable for expression of *patched-2* and recovering the same from the cell culture.

10 In yet another embodiment, the invention provides an isolated *patched-2* polypeptide. In particular, the invention provides isolated native sequence *patched-2* polypeptide, which in one embodiment is a human *patched-2* including an amino acid sequence comprising residues 1 to about 1203 of Figure 1 (SEQ ID NO:2). Human *patched-2* polypeptides with or without the initiating methionine are specifically included. Alternatively, the invention provides a human *patched-2* polypeptide encoded by the nucleic acid deposited under accession number ATCC 209778.

15 In yet another embodiment, the invention provides chimeric molecules comprising a *patched-2* polypeptide fused to a heterologous polypeptide or amino acid sequence. An example of such a chimeric molecule comprises a *patched-2* polypeptide fused to an epitope tag sequence or a constant region of an immunoglobulin.

20 In yet another embodiment, the invention provides expressed sequence tag (EST) comprising the nucleotide sequences identified in Fig. 2A (905531) (SEQ ID NO:3) and Fig. 2B (1326258) (SEQ ID NO:5).

In yet another embodiment, the invention provides for alternatively spliced variants of human *patched-2* having *patched-2* biological activity.

25 In yet another embodiment, the invention provides for method of using *patched-2* for the treatment of disorders which are mediated at least in part by *Hedgehog* (*Hh*), especially *Desert hedgehog* (*Dhh*). In particular, testicular cancer. In yet another embodiment, the invention provides a method of using antagonists or agonists of *patched-2* for treating disorders or creating a desirable physiological condition effected by blocking *Hh* signaling, especially *Dhh* signaling. (E.g, contraception).

BRIEF DESCRIPTION OF THE DRAWINGS

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Figures 1A-1E shows the nucleotide (SEQ ID NO:1) and derived amino acid (SEQ ID NO:2) sequence of a native sequence of human *Ptch-2* polypeptide.

Figure 2A shows EST 905531 (SEQ ID NO:3) and Fig. 2B shows EST 1326258 (SEQ ID NO:5) (SEQ ID NO:4) in alignment with human *Ptch* (SEQ ID NO:18). These ESTs were used in the cloning of human full-length *Ptch-2* (SEQ ID NO:1).

35 Figures 3A-3D show a comparison between human *Ptch* (SEQ ID NO:4) (SEQ ID NO:7) and *Ptch-2* (SEQ ID NO:2). Gaps introduced for optimal alignment are indicated by dashes. Identical amino acids are

boxed. The 12 transmembrane domains are indicated by the gray boxes, all of which are conserved between the two sequences. Alignment results between the two sequences indicate 53% identity. The most significant difference is a shorter C-terminal intracellular domain in human *Ptch-2* (SEQ ID NO:2) in comparison with human *Ptch* (SEQ ID NO:4).

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Figure 4 shows a northern blot of *Ptch-2* (SEQ ID NO:2) which indicates expression is limited to the testis. Multiple human fetal and adult tissue northern blots were probe fragments corresponding to the 3'-untranslated region of murine *Ptch-2*.

Figure 5 shows a chromosomal localization of two BAC clones which were isolated by PCR screening with human *patched-2* derived probes. Both probes were mapped by FISH to human chromosome 1p33-34.

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Figures 6A-F is an *in situ* hybridization comparing *Ptch* (SEQ ID NO:4), *Ptch-2* (SEQ ID NO:2) and *Fused* (FuRK) (SEQ ID NO:10). High magnification of mouse testis showing expression of (A) *Ptch* (SEQ ID NO:4), *Ptch-2* (SEQ ID NO:2) (B) and FuRK (SEQ ID NO:10) (C). Low magnification of testis section hybridized with *Ptch-2* sense (SEQ ID NO:11) (D) and anti-sense probe (SEQ ID NO:12) (E) respectively. Fig. 6(F) shows low magnification of testis section hybridized with FuRK (SEQ ID NO:10). Scale bar: A, B, C: 0.05 mm; D, E, F: 0.33 mm.

Figure 7A is logarithmic plot comparing the binding *Ptch-2* (SEQ ID NO:2) to *Dhh* (SEQ ID NO:13) and *Shh* (SEQ ID NO:14). Competitive binding of recombinant murine ^{125}I -*Shh* to 293 cells overexpressing h*Ptch* (SEQ ID NO:4) or h*Ptch-2* (SEQ ID NO:2). There was no detectable binding to mock transfected cells (data not shown). Figure 7B is a western blot illustrating co-immunoprecipitation of epitope tagged *Ptch* (SEQ ID NO:4) or *Ptch-2* (SEQ ID NO:2) with epitope tagged *Smo* (SEQ ID NO:15). Immunoprecipitation was performed with antibodies to the Flag tagged *Ptch* (SEQ ID NO:4) and analyzed on a 6% acrylamide gel with antibodies to the Myc tagged *Smo* (SEQ ID NO:15). Protein complexes can be detected for both *Ptch* (SEQ ID NO:4) and *Ptch-2* (SEQ ID NO:2) with *Smo* (SEQ ID NO:15). *Ptch* (SEQ ID NO:4) and *Ptch-2* (SEQ ID NO:2) express at similar levels as shown by immunoprecipitation using antibodies to the Flag-tag and western blot using the same anti-Flag antibody.

Figures 8A-8D is a sequence comparison between human *Ptch-2* (SEQ ID NO:2) and murine *Ptch-2* (SEQ ID NO:7) (SEQ ID NO:8), which indicates that there is about 91% identity between the two sequences.

Figure 9 is an *in situ* hybridization which demonstrates the accumulation of *Ptch* (SEQ ID NO:4) and *Ptch-2* (SEQ ID NO:2) mRNA detected by *in situ* hybridization in basal cells of E18 transgenic mice overexpressing *SMO-M2* (SEQ ID NO:16) (Xie *et al.*, *Nature* 391: 90-92 (1998)).

Figures 10A-10D depict a partial sequence representing clone 3A (SEQ ID NO:8) (SEQ ID NO:9), a partial *patched-2* fragment which was initially isolated from a fetal brain library.

Figures 11A-B depict a partial sequence representing clone 16.1 (SEQ ID NO:9) (SEQ ID NO:10), a partial *patched-2* fragment which was isolated from a testis library.

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DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS**I. Definitions**

The terms "*patched-2*" and "*patched-2 polypeptide*" when used herein encompass native sequence *patched-2* and *patched-2* variants (which are further defined herein) having *patched-2* biological activity. 5 *Patched-2* may be isolated from a variety of sources, such as from testes tissue types or from another source, or prepared by recombinant or synthetic methods.

A "native sequence *patched-2*" comprises a polypeptide having the same amino acid sequence as a human *patched-2* derived from nature. Such native sequence *patched-2* can be isolated from nature or can be produced by recombinant and/or synthetic means. The term "native sequence vertebrate *patched-2*" specifically 10 encompasses naturally occurring truncated forms of human *patched-2*, naturally occurring variant forms (e.g., alternatively spliced forms) and naturally-occurring allelic variants of human *patched-2*. Thus, one embodiment of the invention, the native sequence *patched-2* is a mature or full-length native *patched-2* comprising amino acids 1 to 1203 of Fig. 1 (SEQ ID NO:2) with or without the initiating methionine at position 1.

"*Patched-2 variant*" means an active human *patched-2* as defined below having at least > 91% amino 15 acid sequence identity to (a) a DNA molecule encoding a *patched-2* polypeptide, or (b) the complement of the DNA molecule of (a). In a particular embodiment, the *patched-2* variant has at least > 91% amino acid sequence 20 homology with the human *Ptch-2* (SEQ ID NO:2) having the deduced amino acid sequence shown in Fig. 1 for a full-length native sequence human *patched-2*. Such *patched-2* variants include, without limitation, *patched-2* polypeptides wherein one or more amino acid residues are added, or deleted, at the N- or C-terminus of the sequence of Fig. 1 (SEQ ID NO:2). Preferably, the nucleic acid or amino acid sequence identity is at least about 92%, more preferably at least about 93%, and even more preferably at least about 95%.

The term "*Ptch*" or "*Ptch-2*" refer to the particular species of molecules isolated and characterized in the application, while the terms "*patched*" and "*patched-2*" refer to the more generalized description as defined above.

25 "Percent (%) amino acid sequence identity" with respect to the *patched-2* sequences identified herein is defined as the percentage of amino acid residues in a candidate sequence that are identical with the amino acid residues in the *patched-2* sequence, after aligning the sequences and introducing gaps, if necessary, to achieve the maximum percent sequence identity, and not considering any conservative substitutions as part of the sequence identity. Alignment for purposes of determining percent amino acid sequence identity can be achieved 30 in various ways that are within the skill in the art, for instance, using publicly available computer software such as BLAST-2 software that are set to their default parameters. Those skilled in the art can determine appropriate parameters for measuring alignment, including any algorithms needed to achieve maximal alignment over the full length of the sequences being compared. Alternatively, % identity can be determined by Align-2, authored by Genentech, Inc., which was filed with user documentation in the United States Copyright Office, Washington, 35 D.C. 20559, on December 10, 1991, and is registered under U.S. Copyright Registration No. TXU 510087.

"Percent (%) nucleic acid sequence identity" with respect to the *patched-2* sequences identified herein is defined as the percentage of nucleotides in a candidate sequence that are identical with the nucleotides in the

patched-2 sequence, after aligning the sequences and introducing gaps, if necessary, to achieve the maximum percent sequence identity. Alignment for purposes of determining percent nucleic acid sequence identity can be achieved in various ways that are within the skill in the art, for instance, using publicly available computer software such as BLAST-2 software that are set to their default parameters. Those skilled in the art can 5 determine appropriate parameters for measuring alignment, including any algorithms needed to achieve maximal alignment over the full length of the sequences being compared. Alternatively, % identity can be determined by Align-2, authored by Genentech, Inc., which was filed with user documentation in the United States Copyright Office, Washington, D.C. 20559, on December 10, 1991, and is registered under U.S. Copyright Registration No. TXU 510087.

10 The term "epitope tagged" when used herein refers to a chimeric polypeptide comprising *patched-2* polypeptide, or a portion thereof, *patched-2* to a "tag polypeptide". The tag polypeptide has enough residues to provide an epitope against which an antibody can be made, yet is short enough such that it does not interfere with activity of the *patched-2* polypeptide. The tag polypeptide preferably also is fairly unique so that the antibody does not substantially cross-react with other epitopes. Suitable tag polypeptides generally have at least 15 six amino acid residues and usually between about 8 to about 50 amino acid residues (preferably, between about 10 to about 20 residues).

15 As used herein, the term "immunoadhesin" designates antibody-like molecules which combine the binding specificity of a heterologous protein (an "adhesin") with the effector functions of immunoglobulin constant domains. Structurally, the immunoadhesin comprise a fusion of an amino acid sequence with the desired binding specificity which is other than the antigen recognition and binding site of an antibody (*i.e.*, is 20 "heterologous"), and an immunoglobulin constant domain sequence. The adhesin part of an immunoadhesin molecule typically is a contiguous amino acid sequence comprising at least the binding site of a receptor or a ligand. The immunoglobulin constant domain sequence in the immunoadhesins may be obtained from any immunoglobulin, such as IgG-1, IgG-2, IgG-3 or IgG-4 subtypes, IgA (including IgA-1 and IgA-2, IgE, IgD or 25 IgM. Immunoadhesion reported in the literature include fusions of the T cell receptor* [Gascoigne *et al.*, *Proc. Natl. Acad. Sci. USA* 84: 2936-2940 (1987)]; CD4* [Capron *et al.*, *Nature* 337: 525-531 (1989); Traunecker *et al.*, *Nature* 339: 68-70 (1989); Zettmeissl *et al.*, *DNA Cell Biol. USA* 9: 347-353 (1990); Byrn *et al.*, *Nature* 344, 667-670 (1990)]; L-selectin (homing receptor) [Watson *et al.*, *J. Cell. Biol.* 110, 2221-2229 (1990); Watson *et al.*, *Nature* 349, 164-167 (1991)]; CD44* [Aruffo *et al.*, *Cell* 61, 1303-1313 (1990)]; CD28* and B7* [Linsley *et al.*, *J. Exp. Med.* 173, 721-730 (1991)]; CTLA-4* [Lisley *et al.*, *J. Exp. Med.* 174, 561-569 (1991)]; CD22* [Stamenkovic *et al.*, *Cell* 66, 1133-1144 (1991)]; TNF receptor [Ashkenazi *et al.*, *Proc. Natl. Acad. Sci. USA* 88, 10535-10539 30 (1991); Lesslauer *et al.*, *Eur. J. Immunol.* 21, 2883-2886 (1991); Peppel *et al.*, *J. Exp. Med.* 174, 1483-1489 (1991)]; NP receptors [Bennett *et al.*, *J. Biol. Chem.* 266, 23060-23067 (1991)]; IgE receptor α -chain* [Ridgway and Gorman, *J. Cell. Biol.* 115, abstr. 1448 (1991)]; HGF receptor [Mark, M.R. *et al.*, *J. Biol. Chem.*, 267(36): 35 26166-26171 (1992)], where the asterisk (*) indicates that the receptor is a member of the immunoglobulin superfamily.

"Stringency" of hybridization reactions is readily determinable by one of ordinary skill in the art, and generally is an empirical calculation dependent upon probe length, washing temperature, and salt concentration. In general, longer probes require higher temperatures for proper annealing, while shorter probes need lower temperatures. Hybridization generally depends upon the ability of denatured DNA to reanneal when complementary strands are present in an environment near but below their T^m (melting temperature). The higher the degree of desired homology between the probe and hybridizable sequence, the higher the relative temperature which can be used. As a result, it follows that higher relative temperatures would tend to make the reaction conditions more stringent, while lower temperatures less so. Moreover, stringency is also inversely proportional to salt concentrations. For additional details and explanation of stringency of hybridization reactions, see Ausubel *et al.*, *Current Protocols in Molecular Biology* (1995).

"Stringent conditions," as defined herein may be identified by those that: (1) employ low ionic strength and high temperature for washing, for example 0.015 M sodium chloride/0.0015 M sodium citrate/0.1% sodium dodecyl sulfate at 50°C; (2) employ during hybridization a denaturing agent, such as formamide, for example, 50% (vol/vol) formamide with 0.1% bovine serum albumin/0.1% Ficoll/0.1% polyvinylpyrrolidone/50 mM sodium phosphate buffer at pH 6.5 with 750 mM sodium chloride, 75 mM sodium citrate at 42°C; (3) employ 50% formamide, 5 x SSC (0.75 M NaCl, 0.075 M sodium citrate), 50 mM sodium phosphate (pH 6.8), 0.1% sodium pyrophosphate, 5 x Denhardt's solution, sonicated salmon sperm DNA (50 µg/ml), 0.1% SDS, and 10% dextran sulfate at 42°C, with washes at 42°C in 0.2 x SSC (sodium chloride/sodium citrate) and 50% formamide at 55°C, followed by a high-stringency wash consisting of 0.1 x SSC containing EDTA at 55°C.

"Isolated," when used to describe the various polypeptides disclosed herein, means polypeptide that has been identified and separated and/or recovered from a component of its natural environment. Contaminant components of its natural environment are materials that would typically interfere with diagnostic or therapeutic uses for the polypeptide, and may include enzymes, hormones, and other proteinaceous or non-proteinaceous solutes. In preferred embodiments, the polypeptide will be purified (1) to a degree sufficient to obtain at least 15 residues of N-terminal or internal amino acid sequence by use of a spinning cup sequenator, or (2) to homogeneity by SDS-PAGE under non-reducing or reducing conditions using Coomassie blue or, preferably, silver stain. Isolated polypeptide includes polypeptide *in situ* within recombinant cells, since at least one component of the vertebrate *patched-2* natural environment will not be present. Ordinarily, however, isolated polypeptide will be prepared by at least one purification step.

An "isolated" *patched-2* nucleic acid molecule is a nucleic acid molecule that is identified and separated from at least one contaminant nucleic acid molecule with which it is ordinarily associated in the natural source of the *patched-2* nucleic acid. An isolated *patched-2* nucleic acid molecule is other than in the form or setting in which it is found in nature. Isolated *patched-2* nucleic acid molecules therefore are distinguished from the corresponding native *patched-2* nucleic acid molecule as it exists in natural cells.

The term "control sequences" refers to DNA sequences necessary for the expression of an operably linked coding sequence in a particular host organism. The control sequences that are suitable for prokaryotes, for example, include a promoter, optionally an operator sequence, and a ribosome binding site. Eukaryotic cells

are known to utilize promoters, polyadenylation signals, and enhancers.

Nucleic acid is "operably linked" when it is placed into a functional relationship with another nucleic acid sequence. For example, DNA for a presequence or secretory leader is operably linked to DNA for a polypeptide if it is expressed as a preprotein that participates in the secretion of the polypeptide; a promoter or 5 enhancer is operably linked to a coding sequence if it affects the transcription of the sequence; or a ribosome binding site is operably linked to a coding sequence if it is positioned so as to facilitate translation. Generally, "operably linked" means that the DNA sequences being linked are contiguous, and, in the case of a secretory leader, contiguous and in reading frame. However, enhancers do not have to be contiguous. Linking is accomplished by ligation at convenient restriction sites. If such sites do not exist, the synthetic oligonucleotide 10 adaptors or linkers are used in accordance with conventional practice.

The term "antibody" is used in the broadest sense and specifically covers single monoclonal antibodies (including agonist and antagonist antibodies), antibody compositions with polyepitopic specificity, as well as antibody fragments (e.g., Fab, F(ab')₂ and Fv), so long as they exhibit the desired biological activity.

The term "monoclonal antibody" as used herein refers to an antibody obtained from a population of 15 substantially homogeneous antibodies, i.e., the individual antibodies comprising the population are identical except for possible naturally occurring mutations that may be present in minor amounts. Monoclonal antibodies are highly specific, being directed against a single antigenic site. Furthermore, in contrast to conventional (polyclonal) antibody preparations that typically include different antibodies directed against different determinants (epitopes), each monoclonal antibody is directed against a single determinant on the antigen. In 20 addition to their specificity, the monoclonal antibodies are advantageous in that they are synthesized by the hybridoma culture, uncontaminated by other immunoglobulins. The modifier "monoclonal" indicates the character of the antibody as being obtained from a substantially homogeneous population of antibodies, and is not to be construed as requiring production of the antibody by any particular method. For example, the monoclonal antibodies to be used in accordance with the present invention may be made by the hybridoma 25 method first described by Kohler & Milstein, *Nature* 256:495 (1975), or may be made by recombinant DNA methods [see, e.g. U.S. Patent No. 4,816,567 (Cabilly *et al.*)].

The monoclonal antibodies herein specifically include "chimeric" antibodies (immunoglobulins) in which a portion of the heavy and/or light chain is identical with or homologous to corresponding sequences in antibodies derived from a particular species or belonging to a particular antibody class or subclass, while the 30 remainder of the chain(s) is identical with or homologous to corresponding sequences in antibodies derived from another species or belonging to another antibody class or subclass, as well as fragments of such antibodies, so long as they exhibit the desired biological activity [U.S. Patent No. 4,816,567; Cabilly *et al.*; Morrison *et al.*, *Proc. Natl. Acad. Sci. USA* 81, 6851-6855 (1984)].

"Humanized" forms of non-human (e.g. murine) antibodies are chimeric immunoglobulins, 35 immunoglobulin chains or fragments thereof (such as Fv, Fab, Fab', F(ab')₂ or other antigen-binding subsequences of antibodies) which contain minimal sequence derived from non-human immunoglobulin. For the most part, humanized antibodies are human immunoglobulins (recipient antibody) in which residues from a

complementary determining region (CDR) of the recipient are replaced by residues from a CDR of a non-human species (donor antibody) such as mouse, rat or rabbit having the desired specificity, affinity and capacity. In some instances, corresponding non-human residues replace Fv framework residues of the human immunoglobulin. Furthermore, humanized antibody may comprise residues that are found neither in the 5 recipient antibody nor in the imported CDR or framework sequences. These modifications are made to further refine and optimize antibody performance. In general, the humanized antibody will comprise substantially all of at least one, and typically two, variable domains, in which all or substantially all of the CDR regions correspond to those of a non-human immunoglobulin and all or substantially all of the FR regions are those of a human immunoglobulin consensus sequence. The humanized antibody optimally also will comprise at least a portion of 10 an immunoglobulin constant region (Fc), typically that of a human immunoglobulin. For further details see: Jones *et al.*, *Nature* 321, 522-525 (1986); Reichmann *et al.*, *Nature* 332, 323-329 (1988); Presta, *Curr. Op. Struct. Biol.* 2 593-596 (1992) and U.S. Patent No. 5,225,539 (Winter) issued July 6, 1993.

15 "Active" or "activity" for the purposes herein refers to form(s) of *patched-2* which retain the biologic and/or immunologic activities of native or naturally occurring *patched-2*. A preferred activity is the ability to bind to and affect, *e.g.*, block or otherwise modulate, *hedgehog*, especially *Desert hedgehog* signaling. For example, the regulation of the pathogenesis of testicular cancer, male spermatocyte formation and basal cell carcinoma.

20 The term "antagonist" is used herein in the broadest sense to include any molecule which blocks, prevents, inhibits, neutralizes the normal functioning of *patched-2* in the *Hh* signaling pathway. One particular form of antagonist includes a molecule that interferes with the interaction between *Dhh* (SEQ ID NO:13) and *Ptch-2* (SEQ ID NO:2). Alternatively, an antagonist ~~could~~ also be a molecule which increases the levels of 25 *patched-2*. In a similar manner, the term "agonist" is used herein to include any molecule which promotes, enhances or stimulates the binding of a *Hh* to *patched-2* in the *Hh* signaling pathway (*e.g.*, blocking binding of *Ptch-2* (SEQ ID NO:2) to *Smo* (SEQ ID NO:17)). Suitable molecules that affect the protein-protein interaction 30 of *Hh* and *Ptch-2* and its binding proteins include fragments of the latter or small bioorganic molecules, *e.g.*, peptidomimetics, which will prevent or enhance, as the case may be, the binding of *Hh* to *Ptch-2*. Non-limiting examples include proteins, peptides, glycoproteins, glycopeptides, glycolipids, polysaccharides, oligosaccharides, nucleic acids, bioorganic molecules, peptidomimetics, pharmacological agents and their metabolites, transcriptional and translation control sequences, and the like. Another preferred form of antagonist includes antisense oligonucleotides that inhibit proper transcription of wild type *patched-2*.

35 The term "modulation" or "modulating" means upregulation or downregulation of a signaling pathway. Cellular processes under the control of signal transduction may include, but are not limited to, transcription of specific genes; normal cellular functions, such as metabolism, proliferation, differentiation, adhesion, apoptosis and survival, as well as abnormal processes, such as transformation, blocking of differentiation and metastasis.

35 The techniques of "polymerase chain reaction," or "PCR", as used herein generally refers to a procedure wherein minute amounts of a specific piece of nucleic acid, RNA and/or DNA are amplified as described in U.S. Pat. No. 4,683,195 issued 28 July 1987. Generally, sequence information from the ends of the

region of interest or beyond needs to be available, such that oligonucleotide primers can be designed; these primers will be identical or similar in sequence to opposite strands of the template to be amplified. The 5' terminal nucleotides of the two primers may coincide with the ends of the amplified material. PCR sequences form total genomic DNA, and cDNA transcribed from total cellular RNA, bacteriophage, or plasmid sequences, 5 etc. See generally Mullis *et al.*, *Cold Spring Harbor Symp. Quant. Biol.* 51: 263 (1987); Erlich, Ed., PCR Technology, (Stockton Press, NY, 1989). As used herein, PCR is considered to be one, but not the only, example of a nucleic acid test sample comprising the use of a known nucleic acid as a primer and a nucleic acid polymerase to amplify or generate a specific piece of nucleic acid.

10 II. Compositions and Methods of the InventionA. Full-length *patched*-2

The present invention provides newly identified and isolated nucleotide sequences encoding polypeptides referred to in the present application as *patched*-2. In particular, Applicants have identified and isolated cDNA encoding a human *patched*-2 polypeptide, as disclosed in further detail in the Examples below. 15 Using BLAST, BLAST-2 and FastA sequence alignment computer programs (set to the default parameters), Applicants found that a full-length native sequence human *Ptch*-2 (SEQ ID NO:2) (shown in Figure 3) has 53% amino acid sequence identity with human *patched* (SEQ ID NO:4) (SEQ ID NO:7). Moreover human full-length *patched*-2 (SEQ ID NO:2) has about a 91% sequence identity with murine *Ptch*-2 (SEQ ID NO:7) (SEQ ID NO:8) (Fig. 8). Accordingly, it is presently believed that the human *patched*-2 (SEQ ID NO:2) disclosed in the 20 present application is a newly identified member of the mammalian hedgehog signaling cascade, specifically *Desert hedgehog*.

The full-length native sequence of human *patched*-2 gene, or portions thereof, may be used as hybridization probes for a cDNA library to isolate the full-length gene or to isolate still other vertebrate homolog genes (for instance, those encoding naturally-occurring variants of *patched*-2 or *patched*-2 from other species) 25 which have a desired sequence identity to the human *patched*-2 sequence disclosed in Fig.1 (SEQ ID NO:2). Optionally, the length of the probes will be about 20 to about 50 bases. The hybridization probes may be derived from the nucleotide sequence of Fig. 1 (SEQ ID NO:1) or from genomic sequences including promoters, enhancer elements and introns of native sequence vertebrate *patched*-2. By way of example, a screening method 30 will comprise isolating the coding region of the vertebrate *patched*-2 gene using the known DNA sequence to synthesize a selected probe of about 40 bases. Hybridization probes may be labeled by a variety of labels, including radionucleotides such as ³²P or ³⁵S, or enzymatic labels such as alkaline phosphatase coupled to the probe via avidin/biotin coupling systems. Labeled probes having a sequence complementary to that of the vertebrate *patched*-2 gene of the present invention can be used to screen libraries of human cDNA, genomic DNA or mRNA to determine which members of such libraries the probe hybridizes to.

35

B. *Patched*-2 Variants

In addition to the full-length native sequence *patched*-2 described herein, it is contemplated that

patched-2 variants can be prepared. *Patched-2* variants can be prepared by introducing appropriate nucleotide changes into a known *patched-2* DNA, or by synthesis of the desired *patched-2* polypeptides. Those skilled in the art will appreciate that amino acid changes may alter post-translational processes of *patched-2*.

Variations in the native full-length sequence *patched-2* or in various domains of the *patched-2* described herein, can be made, for example, using any of the techniques and guidelines for conservative and non-conservative mutations set forth, for instance, in U.S. Patent No. 5,364,934. Variations may be a substitution, deletion or insertion of one or more codons encoding the *patched-2* that results in a change in the amino acid sequence of *patched-2* as compared with the native sequence *patched-2*. Optionally, the variation is by substitution of at least one amino acid with any other amino acid in one or more of the domains of *patched-2*.

5 Guidance in determining which amino acid residue may be inserted, substituted or deleted without adversely affecting the desired activity may be found by comparing the sequence of the *patched-2* with that of homologous known protein molecules and minimizing the number of amino acid sequence changes made in regions of high homology. Amino acid substitutions can be the result of replacing one amino acid with another amino acid having similar structural and/or chemical properties, such as the replacement of a leucine with a serine, *i.e.*, 10 conservative amino acid replacements. Insertions or deletions may optionally be in the range of 1 to 5 amino acids. The variation allowed may be determined by systematically making insertions, deletions or substitutions of amino acids in the sequence and testing the resulting variants for activity in the *in vitro* assay described in the 15 Examples below.

The variations can be made using methods known in the art such as oligonucleotide-mediated (site-directed) mutagenesis, alanine scanning, and PCR mutagenesis. Site-directed mutagenesis [Carter *et al.*, *Nucl. Acids Res.*, 13:4331 (1986); Zoller *et al.*, *Nucl. Acids Res.*, 10: 6487 (1987)], cassette mutagenesis [Wells *et al.*, *Gene*, 34:315 (1985)], restriction selection mutagenesis [Wells *et al.*, *Philos. Trans. R. Soc. London SerA*, 317:415 (1986)] or other known techniques can be performed on the cloned DNA to produce the vertebrate 20 *patched-2* variant DNA.

Scanning amino acid analysis can also be employed to identify one or more amino acids along a contiguous sequence. Among the preferred scanning amino acids are relatively small, neutral amino acids. Such 25 amino acids include alanine, glycine, serine, and cysteine. Alanine is typically a preferred scanning amino acid among this group because it eliminates the side-chain beyond the beta-carbon and is less likely to alter the main-chain conformation of the variant. Alanine is also typically preferred because it is the most common amino acid.

Further, it is frequently found in both buried and exposed positions [Creighton, *The Proteins*, (W.H. Freeman & Co., N.Y.); Chothia, *J. Mol. Biol.*, 150:1 (1976)]. If alanine substitution does not yield adequate amounts of 30 variant, an isoteric amino acid can be used.

In the comparison between human *Ptch* (SEQ ID NO:4)-(SEQ ID NO:7) and *Ptch-2* (SEQ ID NO:2) sequences depicted in Figure 3, the 12 transmembrane domains are identified in gray, while identical residues 35 are boxed. Gaps are indicated by dashes (-) and are inserted to maximize the total identity score between the two sequences.

C. Modifications of *patched-2*

Covalent modifications of *patched-2* are included within the scope of this invention. One type of covalent modification includes reacting targeted amino acid residues of *patched-2* with an organic derivatizing agent that is capable of reacting with selected side chains or the N- or C-terminal residues of the vertebrate *patched-2*. Derivatization with bifunctional agents is useful, for instance, for crosslinking *patched-2* to a water-insoluble support matrix or surface for use in the method for purifying anti-*patched-2* antibodies, and vice-versa. Commonly used crosslinking agents include, e.g., 1,1-bis(diazo-acetyl)-2-phenylethane, glutaraldehyde, N-hydroxy-succinimide esters, for example, esters with 4-azido-salicylic acid, homobifunctional imidoesters, including disuccinimidyl esters such as 3,3'-dithiobis-(succinimidyl-propionate), bifunctional maleimides such as bis-N-maleimido-1,8-octane and agents such as methyl-3-[(p-azidophenyl)-dithio]propioimidate.

Other modifications include deamidation of glutamyl and asparaginyl residues to the corresponding glutamyl and aspartyl residues, respectively, hydroxylation of proline and lysine, phosphorylation of hydroxyl groups of seryl or threonyl residues, methylation of the α -amino groups of lysine, arginine, and histidine side chains [T.E. Creighton, *Proteins: Structure and Molecular Properties*, W.H. Freeman & Co., San Francisco, pp. 79-86 (1983)], acetylation of the N-terminal amine, and amidation of any C-terminal carboxyl group.

Another type of covalent modification of *patched-2* comprises linking the *patched-2* polypeptide to one of a variety of nonproteinaceous polymers, e.g., polyethylene glycol, polypropylene glycol, or polyoxyalkylenes, in the manner set forth in U.S. Patent Nos. 4,640,835; 4,496,689; 4,301,144; 4,670,417; 4,791,192 or 4,179,337. Such modifications would be expected to increase the half-life of the molecules in circulation in a mammalian system; Extended half-life of *patched-2* molecules might be useful under certain circumstances, such as where the *patched-2* variant is administered as a therapeutic agent.

The *patched-2* of the present invention may also be modified in a way to form a chimeric molecule comprising *patched-2* bonded to another, heterologous polypeptide or amino acid sequence. In one embodiment, such a chimeric molecule comprises a fusion of *patched-2* with a tag polypeptide, which provides an epitope to which an anti-tag antibody can selectively bind. The epitope tag is generally placed at the amino- or carboxyl- terminus of the *patched-2*. The presence of such epitope-tagged forms of the *patched-2* can be detected using an antibody against the tag polypeptide. Also, provision of the epitope tag enables the *patched-2* to be readily purified by affinity purification using an anti-tag antibody or another type of affinity matrix that binds to the epitope tag. In an alternative embodiment, the chimeric molecule may comprise a fusion of the *patched-2* with an immunoglobulin or a particular region of an immunoglobulin. For a bivalent form of the chimeric molecule, such a fusion could be to the Fc region of an IgG molecule. Ordinarily, the C-terminus of a contiguous amino acid sequence of a *patched-2* receptor is fused to the N-terminus of a contiguous amino acid sequence of an immunoglobulin constant region, in place of the variable region(s), however N-terminal fusions are also possible.

Typically, such fusions retain at least functionally active hinge, CH2 and CH3 domains of the constant region of an immunoglobulin heavy chain. Fusions are also made to the C-terminus of the Fc portion of a constant domain, or immediately N-terminal to the CH1 of the heavy chain or the corresponding region of the light chain.

This ordinarily is accomplished by constructing the appropriate DNA sequence and expressing it in recombinant cell culture. Alternatively, immunoadhesins may be synthesized according to known methods.

The precise site at which the fusion is made is not critical; particular sites are well known and may be selected in order to optimize the biological activity, secretion or binding characteristics of the immunoadhesins.

5 In a preferred embodiment, the C-terminus of a contiguous amino acid sequence which comprises the binding site(s) of *patched-2*, at the N-terminal end, to the C-terminal portion of an antibody (in particular the Fc domain), containing the effector functions of an immunoglobulin, *e.g.*, immunoglobulin G₁ (IgG-1). As herein above mentioned, it is possible to fuse the entire heavy chain constant region to the sequence containing the binding site(s). However, more preferably, a sequence beginning in the hinge region just upstream of the papain cleavage 10 site (which defines IgG Fc chemically; residue 216, taking the first residue of heavy chain constant region to be 114 [Kobat *et al.*, *supra*], or analogous sites of other immunoglobulins) is used in the fusion. Although it was earlier thought that in immunoadhesins the immunoglobulin light chain would be required for efficient secretion of the heterologous protein-heavy chain fusion proteins, it has been found that even the immunoadhesins containing the whole IgG1 heavy chain are efficiently secreted in the absence of light chain. Since the light chain is unnecessary, 15 the immunoglobulin heavy chain constant domain sequence used in the construction of the immunoadhesins of the present invention may be devoid of a light chain binding site. This can be achieved by removing or sufficiently altering immunoglobulin heavy chain sequence elements to which the light chain is ordinarily linked so that such binding is no longer possible. Thus, the CH1 domain can be entirely removed in certain embodiments of the *patched-2*/immunoglobulin chimeras.

20 In a particularly preferred embodiment, the amino acid sequence containing the extracellular domain(s) of *patched-2* is fused to the hinge region and CH2, CH3; or CH1, hinge, CH2 and CH3 domains of an IgG-1, IgG-2, IgG-3, or IgG-4 heavy chain.

25 In some embodiments, the *patched-2*/immunoglobulin molecules (immunoadhesins) are assembled as monomers, dimers or multimers, and particularly as dimers or tetramers. Generally, these assembled immunoadhesins will have known unit structures similar to those of the corresponding immunoglobulins. A basic four chain structural unit (a dimer of two immunoglobulin heavy chain-light chain pairs) is the form in which IgG, IgA and IgE exist. A four chain unit is repeated in the high molecular weight immunoglobulins; IgM generally exists as a pentamer of basic four-chain units held together by disulfide bonds. IgA globulin, and occasionally IgG globulin, may also exist in a multimeric form in serum. In the case of multimers, each four chain unit may be the 30 same or different.

It is not necessary that the entire immunoglobulin portion of the *patched-2*/immunoglobulin chimeras be from the same immunoglobulin. Various portions of different immunoglobulins may be combined, and variants and derivatives of native immunoglobulins can be made as herein above described with respect to *patched-2*, in order to optimize the properties of the immunoadhesin molecules. For example, immunoadhesin constructs in which the hinge of IgG-1 was replaced with that of IgG-3 were found to be functional and showed pharmacokinetics 35 comparable to those of immunoadhesins comprising the entire IgG-1 heavy chain.

Various tag polypeptides and their respective antibodies are well known in the art. Examples include poly-histidine (poly-his) or poly-histidine-glycine (poly-his-gly) tags; the flu HA tag polypeptide and its antibody 12CA5 [Field *et al.*, *Mol. Cell. Biol.*, 8: 2159-2165 (1988)]; the c-myc tag and the 8F9, 3C7, 6E10, G4, B7 and 9E10 antibodies thereto [Evan *et al.*, *Molecular and Cellular Biology*, 5:3610-3616 (1985)]; and the Herpes Simplex virus glycoprotein D (gD) tag and its antibody [Paborsky *et al.*, *Protein Engineering*, 3(6):547-553 (1990)]. Other tag polypeptides include the Flag-peptide [Hopp *et al.*, *BioTechnology*, 6:1204-1210 (1988)]; the KT3 epitope peptide [Martin *et al.*, *Science*, 255:192-194 (1992)]; an α -tubulin epitope peptide [Skinner *et al.*, *J. Biol. Chem.*, 266:15163-15166 (1991)]; and the T7 gene 10 protein peptide tag [Lutz-Freyermuth *et al.*, *Proc. Natl. Acad. Sci. USA*, 87:6393-6397 (1990)]. A preferred tag is the influenza HA tag.

10

D. Preparation of *patched-2*

The description below relates primarily to production of a particular *patched-2* by culturing cells transformed or transfected with a vector containing *patched-2* nucleic acid. It is, of course, contemplated that alternative methods, which are well known in the art, may be employed to prepare *patched-2*. For instance, the *patched-2* sequence, or portions thereof, may be produced by direct peptide synthesis using solid-phase techniques [see, *e.g.*, Stewart *et al.*, *Solid-Phase Peptide Synthesis*, W.H. Freeman Co., San Francisco, CA (1969); Merrifield, *J. Am. Chem. Soc.*, 85:2149-2154 (1963)]. *In vitro* protein synthesis may be performed using manual techniques or by automation. Automated synthesis may be accomplished, for instance, using an Applied Biosystems Peptide Synthesizer (Foster City, CA) using manufacturer's instructions. Various portions 20 of the vertebrate *patched-2* may be chemically synthesized separately and combined using chemical or enzymatic methods to produce the full-length *patched-2*.

1. Isolation of DNA encoding vertebrate *patched-2*

DNA encoding *patched-2* may be obtained from a cDNA library prepared from tissue believed to possess the *patched-2* mRNA and to express it at a detectable level. Accordingly, human *patched-2* DNA can be conveniently obtained from a cDNA library prepared from human tissue, such as described in the Examples. The vertebrate *patched-2*-encoding gene may also be obtained from a genomic library or by oligonucleotide synthesis.

Libraries can be screened with probes (such as antibodies to the *patched-2* or oligonucleotides of at least about 20-80 bases) designed to identify the gene of interest or the protein encoded by it. Screening the cDNA or genomic library with the selected probe may be conducted using standard procedures, such as described in Sambrook *et al.*, *Molecular Cloning: A Laboratory Manual* (New York: Cold Spring Harbor Laboratory Press, 1989). An alternative means to isolate the gene encoding vertebrate *patched-2* is to use PCR methodology [Sambrook *et al.*, *supra*; Dieffenbach *et al.*, *PCR Primer: A Laboratory Manual* (Cold Spring Harbor Laboratory Press, 1995)].

The Examples below describe techniques for screening a cDNA library. The oligonucleotide sequences selected as probes should be of sufficient length and sufficiently unambiguous that false positives are minimized.

The oligonucleotide is preferably labeled such that it can be detected upon hybridization to DNA in the library being screened. Methods of labeling are well known in the art, and include the use of radiolabels like ^{32}P -labeled ATP, biotinylation or enzyme labeling. Hybridization conditions, including moderate stringency and high stringency, are provided in Sambrook *et al.*, *supra*.

5 Sequences identified in such library screening methods can be compared and aligned to other known sequences deposited and available in public databases such as GenBank or other private sequence databases. Sequence identity (at either the amino acid or nucleotide level) within defined regions of the molecule or across the full-length sequence can be determined through sequence alignment using computer software programs such as BLAST, BLAST-2, ALIGN, DNAstar, and INHERIT which employ various algorithms to measure 10 homology.

Nucleic acid having protein coding sequence may be obtained by screening selected cDNA or genomic libraries using the deduced amino acid sequence disclosed herein for the first time, and, if necessary, using conventional primer extension procedures as described in Sambrook *et al.*, *supra*, to detect precursors and processing intermediates of mRNA that may not have been reverse-transcribed into cDNA.

15

2. Selection and Transformation of Host Cells

Host cells are transfected or transformed with expression or cloning vectors described herein for *patched-2* production and cultured in conventional nutrient media modified as appropriate for inducing 20 promoters, selecting transformants, or amplifying the genes encoding the desired sequences. The culture conditions, such as media, temperature, pH and the like, can be selected by the skilled artisan without undue experimentation. In general, principles, protocols, and practical techniques for maximizing the productivity of cell cultures can be found in *Mammalian Cell Biotechnology: A Practical Approach*, M. Butler, ed. (IRL Press, 1991) and Sambrook *et al.*, *supra*.

Methods of transfection are known to the ordinarily skilled artisan, for example, CaPO_4 and 25 electroporation. Depending on the host cell used, transformation is performed using standard techniques appropriate to such cells. The calcium treatment employing calcium chloride, as described in Sambrook *et al.*, *supra*, or electroporation is generally used for prokaryotes or other cells that contain substantial cell-wall barriers. Infection with *Agrobacterium tumefaciens* is used for transformation of certain plant cells, as described by Shaw *et al.*, *Gene*, 23:315 (1983) and WO 89/05859 published 29 June 1989. For mammalian cells without 30 such cell walls, the calcium phosphate precipitation method of Graham and van der Eb, *Virology* 52:456-457 (1978) can be employed. General aspects of mammalian cell host system transformations have been described in U.S. Patent No. 4,399,216. Transformations into yeast are typically carried out according to the method of Van Solingen *et al.*, *J. Bact.*, 130:946 (1977) and Hsiao *et al.*, *Proc. Natl. Acad. Sci. (USA)*, 76:3829 (1979). However, other methods for introducing DNA into cells, such as by nuclear microinjection, electroporation, 35 bacterial protoplast fusion with intact cells, or polycations, *e.g.*, polybrene, polyornithine, may also be used. For various techniques for transforming mammalian cells, see Keown *et al.*, *Methods in Enzymology*, 185:527-537 (1990) and Mansour *et al.*, *Nature*, 336:348-352 (1988).

Suitable host cells for cloning or expressing the DNA in the vectors herein include prokaryote, yeast, or higher eukaryote cells. Suitable prokaryotes include but are not limited to eubacteria, such as Gram-negative or Gram-positive organisms, for example, Enterobacteriae such as *E. coli*. Various *E. coli* strains are publicly available, such as *E. coli* K12 strain MM294 (ATCC 31,446); *E. coli* X1776 (ATCC 31,537); *E. coli* strain 5 W3110 (ATCC 27,325) and K5 772 (ATCC 53,635).

In addition to prokaryotes, eukaryotic microbes such as filamentous fungi or yeast are suitable cloning or expression hosts for vertebrate *patched-2*-encoding vectors. *Saccharomyces cerevisiae* is a commonly used lower eukaryotic host microorganism.

Suitable host cells for the expression of vertebrate *patched-2* are derived from multicellular organisms. 10 Examples of invertebrate cells include insect cells such as *Drosophila* S2 and *Spodoptera* Sf9, as well as plant cells. Examples of useful mammalian host cell lines include Chinese hamster ovary (CHO) and COS cells. More specific examples include monkey kidney CV1 line transformed by SV40 (COS-7, ATCC CRL 1651); human embryonic kidney line (293 or 293 cells subcloned for growth in suspension culture, Graham *et al.*, *J. Gen Virol.*, 36:59 (1977)); Chinese hamster ovary cells/-DHFR (CHO, Urlaub and Chasin, *Proc. Natl. Acad. Sci. USA*, 77:4216 (1980)); mouse sertoli cells (TM4, Mather, *Biol. Reprod.* 23:243-251 (1980)); human lung cells 15 (W138, ATCC CCL 75); human liver cells (Hep G2, HB 8065); and mouse mammary tumor (MMT 060562, ATCC CCL51). The selection of the appropriate host cell is deemed to be within the skill in the art.

3. Selection and Use of a Replicable Vector

20 The nucleic acid (e.g., cDNA or genomic DNA) encoding *patched-2* may be inserted into a replicable vector for cloning (amplification of the DNA) or for expression. Various vectors are publicly available. The vector may, for example, be in the form of a plasmid, cosmid, viral particle, or phage. The appropriate nucleic acid sequence may be inserted into the vector by a variety of procedures. In general, DNA is inserted into an appropriate restriction endonuclease site(s) using techniques known in the art. Vector components generally 25 include, but are not limited to, one or more of a signal sequence, an origin of replication, one or more marker genes, an enhancer element, a promoter, and a transcription termination sequence. Construction of suitable vectors containing one or more of these components employs standard ligation techniques, which are known to the skilled artisan.

Both expression and cloning vectors contain a nucleic acid sequence that enables the vector to replicate 30 in one or more selected host cells. Such sequences are well known for a variety of bacteria, yeast, and viruses. The origin of replication from the plasmid pBR322 is suitable for most Gram-negative bacteria, the 2 μ plasmid origin is suitable for yeast, and various viral origins (SV40, polyoma, adenovirus, VSV or BPV) are useful for cloning vectors in mammalian cells. A preferred replicable expression vector is the plasmid pRK5. *Holmes et al.*, *Science*, 253:1278-1280 (1991).

35 Expression and cloning vectors will typically contain a selection gene, also termed a selectable marker. Typical selection genes encode proteins that (a) confer resistance to antibiotics or other toxins, e.g., ampicillin, neomycin, methotrexate, or tetracycline, (b) complement auxotrophic deficiencies, or (c) supply critical nutrients

not available from complex media, *e.g.*, the gene encoding D-alanine racemase for *Bacilli*.

An example of suitable selectable markers for mammalian cells are those that enable the identification of cells competent to take up the *patched-2* nucleic acid, such as DHFR or thymidine kinase. An appropriate host cell when wild-type DHFR is employed is the CHO cell line deficient in DHFR activity, prepared and propagated as described by Urlaub *et al.*, *Proc. Natl. Acad. Sci. USA*, 77:4216 (1980). A suitable selection gene for use in yeast is the *trp1* gene present in the yeast plasmid YRp7 [Stinchcomb *et al.*, *Nature*, 282:39 (1979); Kingsman *et al.*, *Gene*, 7:141 (1979); Tschemper *et al.*, *Gene*, 10:157 (1980)]. The *trp1* gene provides a selection marker for a mutant strain of yeast lacking the ability to grow in tryptophan, for example, ATCC No. 44076 or PEP4-1 [Jones, *Genetics*, 85:12 (1977)].

Expression and cloning vectors usually contain a promoter operably linked to the *patched-2* nucleic acid sequence to direct mRNA synthesis. Promoters recognized by a variety of potential host cells are well known. Promoters suitable for use with prokaryotic hosts include the β -lactamase and lactose promoter systems [Chang *et al.*, *Nature*, 275:615 (1978); Goeddel *et al.*, *Nature*, 281:544 (1979)], alkaline phosphatase, a tryptophan (*trp*) promoter system [Goeddel, *Nucleic Acids Res.*, 8:4057 (1980); EP 36,776], and hybrid promoters such as the tac promoter [deBoer *et al.*, *Proc. Natl. Acad. Sci. USA*, 80:21-25 (1983)]. Promoters for use in bacterial systems also will contain a Shine-Dalgarno (S.D.) sequence operably linked to the DNA encoding *patched-2*.

Examples of suitable promoting sequences for use with yeast hosts include the promoters for 3-phosphoglycerate kinase [Hitzeman *et al.*, *J. Biol. Chem.*, 255:2073 (1980)] or other glycolytic enzymes [Hess *et al.*, *J. Adv. Enzyme Reg.*, 7:149 (1968); Holland, *Biochemistry*, 17:4900 (1978)], such as enolase, glyceraldehyde-3-phosphate dehydrogenase, hexokinase, pyruvate decarboxylase, phospho-fructokinase, glucose-6-phosphate isomerase, 3-phosphoglycerate mutase, pyruvate kinase, triosephosphate isomerase, phosphoglucose isomerase, and glucokinase.

Other yeast promoters, which are inducible promoters having the additional advantage of transcription controlled by growth conditions, are the promoter regions for alcohol dehydrogenase 2, isocytchrome C, acid phosphatase, degradative enzymes associated with nitrogen metabolism, metallothionein, glyceraldehyde-3-phosphate dehydrogenase, and enzymes responsible for maltose and galactose utilization. Suitable vectors and promoters for use in yeast expression are further described in EP 73,657.

Patched-2 transcription from vectors in mammalian host cells is controlled, for example, by promoters obtained from the genomes of viruses such as polyoma virus, fowlpox virus (UK 2,211,504 published 5 July 1989), adenovirus (such as Adenovirus 2), bovine papilloma virus, avian sarcoma virus, cytomegalovirus, a retrovirus, hepatitis-B virus and Simian Virus 40 (SV40), from heterologous mammalian promoters, *e.g.*, the actin promoter or an immunoglobulin promoter, and from heat-shock promoters, provided such promoters are compatible with the host cell systems.

Inserting an enhancer sequence into the vector may increase transcription of a DNA encoding the vertebrate *patched-2* by higher eukaryotes. Enhancers are cis-acting elements of DNA, usually about from 10 to 300 bp, that act on a promoter to increase its transcription. Many enhancer sequences are now known from

mammalian genes (globin, elastase, albumin, α -fetoprotein, and insulin). Typically, however, one will use an enhancer from a eukaryotic cell virus. Examples include the SV40 enhancer on the late side of the replication origin (bp 100-270), the cytomegalovirus early promoter enhancer, the polyoma enhancer on the late side of the replication origin, and adenovirus enhancers. The enhancer may be spliced into the vector at a position 5' or 3' to the *patched-2* coding sequence, but is preferably located at a site 5' from the promoter.

5 Expression vectors used in eukaryotic host cells (yeast, fungi, insect, plant, animal, human, or nucleated cells from other multicellular organisms) will also contain sequences necessary for the termination of transcription and for stabilizing the mRNA. Such sequences are commonly available from the 5' and, occasionally 3', untranslated regions of eukaryotic or viral DNAs or cDNAs. These regions contain nucleotide 10 segments transcribed as polyadenylated fragments in the untranslated portion of the mRNA encoding *patched-2*.

Still other methods, vectors, and host cells suitable for adaptation to the synthesis of *patched-2* in recombinant vertebrate cell culture are described in Gething *et al.*, *Nature*, 293:620-625 (1981); Mantei *et al.*, *Nature*, 281:40-46 (1979); EP 117,060; and EP 117,058.

15

4. Detecting Gene Amplification/Expression

Gene amplification and/or expression may be measured in a sample directly, for example, by conventional Southern blotting, Northern blotting to quantitate the transcription of mRNA [Thomas, *Proc. Natl. Acad. Sci. USA*, 77:5201-5205 (1980)], dot blotting (DNA analysis), or *in situ* hybridization, using an 20 appropriately labeled probe, based on the sequences provided herein. Alternatively, antibodies may be employed that can recognize specific duplexes, including DNA duplexes, RNA duplexes, and DNA-RNA hybrid duplexes or DNA-protein duplexes. The antibodies in turn may be labeled and the assay may be carried out where the duplex is bound to a surface, so that upon the formation of duplex on the surface, the presence of antibody bound to the duplex can be detected.

25

Gene expression, alternatively, may be measured by immunological methods, such as immunohistochemical staining of cells or tissue sections and assay of cell culture or body fluids, to quantitate directly the expression of gene product. Antibodies useful for immunohistochemical staining and/or assay of sample fluids may be either monoclonal or polyclonal, and may be prepared in any mammal. Conveniently, the 30 antibodies may be prepared against a native sequence *patched-2* polypeptide or against a synthetic peptide based on the DNA sequences provided herein or against exogenous sequence *patched-2* to *patched-2* DNA and encoding a specific antibody epitope.

5. Purification of Polypeptide

Forms of *patched-2* may be recovered from host cell lysates. Since *patched-2* is membrane-bound, it 35 can be released from the membrane using a suitable detergent solution (e.g. Triton-X 100) or by enzymatic cleavage. Cells employed in expression of *patched-2* can be disrupted by various physical or chemical means, such as freeze-thaw cycling, sonication, mechanical disruption, or cell lysing agents.

It may be desired to purify *patched-2* from recombinant cell proteins or polypeptides. The following procedures are exemplary of suitable purification procedures: by fractionation on an ion-exchange column; ethanol precipitation; reverse phase HPLC; chromatography on silica or on a cation-exchange resin such as DEAE; chromatofocusing; SDS-PAGE; ammonium sulfate precipitation; gel filtration using, for example, 5 Sephadex G-75; protein A Sepharose columns to remove contaminants such as IgG; and metal chelating columns to bind epitope-tagged forms of the *patched-2*. Various methods of protein purification may be employed and such methods are known in the art and described for example in Deutscher, *Methods in Enzymology*, 182 (1990); Scopes, *Protein Purification: Principles and Practice*, Springer-Verlag, New York (1982). The purification 10 step(s) selected will depend, for example, on the nature of the production process used and the particular *patched-2* produced.

E. Uses for *patched-2*

(1) *Patched-2 is a specific receptor for Dhh*

The *hedgehog* signaling pathway has been implicated in the formation of embryonic structures in 15 mammals and invertebrates. The multi-pass transmembrane receptor *Ptch*, is a negative regulator of the hedgehog pathway, repressing the serpentine signaling molecule *Smoothened (Smo)*. Data have shown that loss of *Ptch* leads to deregulation of the hedgehog pathway leading to formation of aberrant structures in the embryos and carcinoma in the adult.

Applicants' newly identified second human *patched* gene, termed *patched-2*, has a similar 12 20 transmembrane domain topology as does *patched*, and can bind to all the members of the *Hh* family and can complex with *Smo*-(SEQ ID NO:17). However, the expression patterns of *Ptch-2* and *Ptch* do not overlap. *Ptch-2* is expressed mainly in the developing spermatocytes, which are supported directly by the *Dhh* producing Sertoli cells, which suggests that *Ptch-2* is a receptor for Desert hedgehog.

In the adult tubule, Sertoli cells, which are unusually large secretory cells, traverse the seminiferous 25 tubule from the basal lamina to the luminal aspect, sending out cytoplasmic protrusions that engulf the germ cells. These contacts are particularly close during spermiogenesis, in which the haploid round spermatids undergo differentiation to produce the highly specialized, motile sperm. Tight junctions between adjacent Sertoli cells compartmentalize the tubule into a basal region, which contains mitotic spermatogonia and early spermatocytes, and an adluminal compartment, which contains meiotic spermatocytes and maturing spermatids. In 30 fact, a Sertoli-derived cell line supports the meiotic progression of germ cells in culture, consistent with the view that factors derived from Sertoli cells contribute to germ cell maturation, Rassoulzadegan, M., *et al.*, *Cell* 1993, 75: 997-1006. Loss of *Dhh* activity results in a recessive, sex-specific phenotype. Female mice homozygous for the mutation were fully viable and fertile, whereas male mice were viable but infertile. A gross examination indicated that, as early as 18.5 dpc, the testes of mutant males were noticeably smaller than those of 35 heterozygous littermates. Bitgood *et al.*, *Curr. Biol.*, 1996 6(3): 298-304. Thus, Sertoli cells likely independently regulate mitotic and meiotic stages of germ cell development during postnatal development. Therefore, since *patched-2* appears to be the receptor for *Dhh*-(SEQ ID NO:13), molecules which modulate the

binding of *Dhh* (SEQ ID NO:13) to *Ptch-2* would affect the activation of *Dhh* signaling, and thereby would have utility in the treatment of conditions which are modulated by *Dhh* (SEQ ID NO:13). (For example, testicular cancer). Alternatively, it is also provided that antagonists or agonists of *patched-2* may be used for treating disorders or creating a desirable physiological condition effected by blocking *Dhh* (SEQ ID NO:13) signaling. (E.g., contraception, infertility treatment).

5

(2) *General uses for patched-2*

Nucleotide sequences (or their complement) encoding *patched-2* have various applications in the art of molecular biology, including uses as hybridization probes, in chromosome and gene mapping and in the 10 generation of anti-sense RNA and DNA. *Patched-2* nucleic acid will also be useful for the preparation of *patched-2* polypeptides by the recombinant techniques described herein.

The full-length native sequence *patched-2* gene, or portions thereof, may be used as hybridization probes for a cDNA library to isolate the full-length gene or to isolate still other genes (for instance, those encoding naturally-occurring variants of *patched-2*) which have a desired sequence identity to the *patched-2* 15 sequence disclosed in Fig. 1 (SEQ ID NO:1). Optionally, the length of the probes will be about 20 to about 50 bases. The hybridization probes may be derived from the nucleotide sequence of Fig. 1 (SEQ ID NO:1) or from genomic sequences including promoters, enhancer elements and introns of native sequence *patched-2*. By way of example, a screening method will comprise isolating the coding region of the *patched-2* gene using the known 20 DNA sequence to synthesize a selected probe of about 40 bases. Hybridization probes may be labeled by a variety of labels, including radionucleotides such as ^{32}P or ^{35}S , or enzymatic labels such as alkaline phosphatase coupled to the probe via avidin/biotin coupling systems. Labeled probes having a sequence complementary to that of the *patched-2* gene of the present invention can be used to screen libraries of human cDNA, genomic DNA or mRNA to determine to which members of such libraries the probe hybridizes. Hybridization techniques are described in further detail in the Examples below.

25

The probes may also be employed in PCR techniques to generate a pool of sequences for identification of closely related *patched-2* sequences.

Nucleotide sequences encoding *patched-2* can also be used to construct hybridization probes for mapping the gene, which encodes *patched-2* and for the genetic analysis of individuals with genetic disorders. The nucleotide sequences provided herein may be mapped to a chromosome and specific regions of a 30 chromosome using known techniques, such as *in situ* hybridization, linkage analysis against known chromosomal markers, and hybridization screening with libraries.

35

Patched-2 polypeptides can be used in assays to identify the other proteins or molecules involved in complexing with *patched-2* which ultimately results in the modulation of hedgehog signaling. Alternatively, these molecules can modulate the binding of *patched-2* to *Dhh* (SEQ ID NO:13). By such methods, inhibitors of the binding interaction can be identified. Proteins involved in such binding interactions can also be used to screen for peptide or small molecule inhibitors or agonists of the binding interaction. Also, the substrate of *patched-2* can be used to isolate correlative complexing proteins. Screening assays can be designed to find lead

compounds that mimic the biological activity of a native *patched-2* or to find those that act as a substrate for *patched-2*. Such screening assays will include assays amenable to high-throughput screening of chemical libraries, making them particularly suitable for identifying small molecule drug candidates. Such small molecule inhibitors could block the enzymatic action of *patched-2*, and thereby inhibit *hedgehog* signaling. Small 5 molecules contemplated include synthetic organic or inorganic compounds. The assays can be performed in a variety of formats, including protein-protein binding assays, biochemical screening assays, immunoassays and cell based assays, which are well characterized in the art.

Nucleic acids which encode *patched-2* or its modified forms can also be used to generate either transgenic animals or "knock out" animals which, in turn, are useful in the development and screening of 10 therapeutically useful reagents. A transgenic animal (e.g., a mouse or rat) is an animal having cells that contain a transgene, was introduced into the animal or an ancestor of the animal at a prenatal, e.g., an embryonic stage. A transgene is a DNA sequence that is integrated into the genome of a cell from which a transgenic animal develops. In one embodiment, cDNA encoding *patched-2* can be used to clone genomic DNA encoding 15 *patched-2* in accordance with established techniques and the genomic sequences used to generate transgenic animals that contain cells which express DNA encoding *patched-2*. Methods for generating transgenic animals, particularly animals such as mice or rats, have become conventional in the art and are described, for example, in U.S. Patent Nos. 4,736,866 and 4,870,009. Typically, particular cells would be targeted for *patched-2* transgene incorporation with tissue-specific enhancers. Transgenic animals that include a copy of a transgene encoding *patched-2* introduced into the germ line of the animal at an embryonic stage can be used to examine the effect of 20 increased expression of DNA encoding *patched-2*. Such animals can be used as tester animals for reagents thought to confer protection from, for example, pathological conditions associated with its overexpression.

Non-human homologues of vertebrate *patched-2* can be used to construct a *patched-2* "knock out" animal which has a defective or altered gene encoding *patched-2* as a result of homologous recombination between the endogenous gene encoding *patched-2* and altered genomic DNA encoding *patched-2* introduced 25 into an embryonic cell of the animal. For example, cDNA encoding *patched-2* can be used to clone genomic DNA encoding *patched-2* in accordance with established techniques. A portion of the genomic DNA encoding *patched-2* can be deleted or replaced with another gene, such as a gene encoding a selectable marker that can be used to monitor integration. Typically, several kilobases of unaltered flanking DNA (both at the 5' and 3' ends) 30 are included in the vector [see e.g., Thomas and Capecchi, *Cell*, 51:503 (1987) for a description of homologous recombination vectors]. The vector is introduced into an embryonic stem cell line (e.g., by electroporation) and cells in which the introduced DNA has homologously recombined with the endogenous DNA are selected [see e.g., Li *et al.*, *Cell*, 69:915 (1992)]. The selected cells are then injected into a blastocyst of an animal (e.g., a mouse or rat) to form aggregation chimeras [see e.g., Bradley, in *Teratocarcinomas and Embryonic Stem Cells: A Practical Approach*, E. J. Robertson, ed. (IRL, Oxford, 1987), pp. 113-152]. A chimeric embryo can then be 35 implanted into a suitable pseudopregnant female foster animal and the embryo brought to term to create a "knock out" animal. Progeny harboring the homologously recombined DNA in their germ cells can be identified by standard techniques and used to breed animals in which all cells of the animal contain the homologously

recombined DNA. Knockout animals can be characterized for instance, for their ability to defend against certain pathological conditions and for their development of pathological conditions due to absence of the *patched-2* polypeptide.

Suppression or inhibition (antagonism) of *Dhh* signaling is also an objective of therapeutic strategies.

5 Since *patched-2* can combine with all members of the hedgehog family (i.e., *Shh*, *Dhh*, *Ihh*), antagonist molecules which prevent the binding of hedgehog molecules to *Ptch-2* (SEQ ID NO:2) have therapeutic utility. For example, *SHh* signaling is known to be activated in Basal Cell Carcinoma; *Dhh* (SEQ ID NO:13) is known to be involved in the regulation of spermatogenesis. Inhibitor or antagonist of *Hh* signaling would be effective therapeutics in the treatment of Basal Cell Carcinoma or male contraception, respectively.

10 The stimulation of *Dhh* signaling (agonism) is also an objective of therapeutic strategies. Since *Ptch-2* (SEQ ID NO:2) also binds to the other members of the *Hh* family, *Ihh* and *Shh*, activating *Dhh* signaling would be useful in disease states or disorders characterized by inactive or insufficient *Hh* signaling. For example, degenerative disorders of the nervous system, e.g., Parkinson's disease, memory deficits, Alzheimer's disease, Lou Gehrig's disease, Huntington's disease, schizophrenia, stroke and drug addiction. Additionally, *patched-2* agonists could be used to treat gut diseases, bone diseases, skin diseases, diseases of the testis (including 15 infertility), ulcers, lung diseases, diseases of the pancreas, diabetes, osteoporosis.

F. Anti-patched-2 Antibodies

The present invention further provides anti- vertebrate *patched-2* antibodies. Exemplary antibodies 20 include polyclonal, monoclonal, humanized, bispecific, and heteroconjugate antibodies.

1. Polyclonal Antibodies

The anti-*patched-2* antibodies may comprise polyclonal antibodies. Methods of preparing polyclonal antibodies are known to the skilled artisan. Polyclonal antibodies can be raised in a mammal, for example, by 25 one or more injections of an immunizing agent and, if desired, an adjuvant. Typically, the immunizing agent and/or adjuvant will be injected in the mammal by multiple subcutaneous or intraperitoneal injections. The immunizing agent may include the *patched-2* polypeptide or a fusion protein thereof. It may be useful to conjugate the immunizing agent to a protein known to be immunogenic in the mammal being immunized. Examples of such immunogenic proteins include but are not limited to keyhole limpet hemocyanin, serum 30 albumin, bovine thyroglobulin, and soybean trypsin inhibitor. Examples of adjuvants that may be employed include Freund's complete adjuvant and MPL-TDM adjuvant (monophosphoryl Lipid A, synthetic trehalose dicorynomycolate). The immunization protocol may be selected by one skilled in the art without undue experimentation.

35 2. Monoclonal Antibodies

The anti-*patched-2* antibodies may, alternatively, be monoclonal antibodies. Monoclonal antibodies may be prepared using hybridoma methods, such as those described by Kohler and Milstein, *Nature*, 256:495

(1975). In a hybridoma method, a mouse, hamster, or other appropriate host animal, is typically immunized with an immunizing agent to elicit lymphocytes that produce or are capable of producing antibodies that will specifically bind to the immunizing agent. Alternatively, the lymphocytes may be immunized *in vitro*.

The immunizing agent will typically include the *patched-2* polypeptide or a fusion protein thereof. 5 Generally, either peripheral blood lymphocytes ("PBLs") are used if cells of human origin are desired, or spleen cells or lymph node cells are used if non-human mammalian sources are desired. The lymphocytes are then fused with an immortalized cell line using a suitable fusing agent, such as polyethylene glycol, to form a hybridoma cell [Goding, *Monoclonal Antibodies: Principles and Practice*, Academic Press, (1986) pp. 59-103]. Immortalized cell lines are usually transformed mammalian cells, particularly myeloma cells of rodent, bovine 10 and human origin. Usually, rat or mouse myeloma cell lines are employed. The hybridoma cells may be cultured in a suitable culture medium that preferably contains one or more substances that inhibit the growth or survival of the unfused immortalized cells. For example, if the parental cells lack the enzyme hypoxanthine guanine phosphoribosyl transferase (HPRT or HPRT), the culture medium for the hybridomas typically will include hypoxanthine, aminopterin, and thymidine ("HAT medium"), which substances prevent the growth of 15 HGPRT-deficient cells.

Preferred immortalized cell lines are those that fuse efficiently, support stable high level expression of antibody by the selected antibody-producing cells, and are sensitive to a medium such as HAT medium. More preferred immortalized cell lines are murine myeloma lines, which can be obtained, for instance, from the Salk Institute Cell Distribution Center, San Diego, California and the American Type Culture Collection, Rockville, 20 Maryland. Human myeloma and mouse-human heteromyeloma cell lines also have been described for the production of human monoclonal antibodies [Kozbor, *J. Immunol.*, 133:3001 (1984); Brodeur *et al.*, *Monoclonal Antibody Production Techniques and Applications*, Marcel Dekker, Inc., New York, (1987) pp. 51-63].

The culture medium in which the hybridoma cells are cultured can then be assayed for the presence of 25 monoclonal antibodies directed against *patched-2*. Preferably, the binding specificity of monoclonal antibodies produced by the hybridoma cells is determined by immunoprecipitation or by an *in vitro* binding assay, such as radioimmunoassay (RIA) or enzyme-linked immunoabsorbent assay (ELISA). Such techniques and assays are known in the art. The binding affinity of the monoclonal antibody can, for example, be determined by the Scatchard analysis of Munson and Pollard, *Anal. Biochem.*, 107:220 (1980).

30 After the desired hybridoma cells are identified, the clones may be subcloned by limiting dilution procedures and grown by standard methods [Goding, *supra*]. Suitable culture media for this purpose include, for example, Dulbecco's Modified Eagle's Medium and RPMI-1640 medium. Alternatively, the hybridoma cells may be grown *in vivo* as ascites in a mammal.

The monoclonal antibodies secreted by the subclones may be isolated or purified from the culture 35 medium or ascites fluid by conventional immunoglobulin purification procedures such as, for example, protein A-Sepharose, hydroxylapatite chromatography, gel electrophoresis, dialysis, or affinity chromatography.

The monoclonal antibodies may also be made by recombinant DNA methods, such as those described

in U.S. Patent No. 4,816,567. DNA encoding the monoclonal antibodies of the invention can be readily isolated and sequenced using conventional procedures (e.g., by using oligonucleotide probes that are capable of binding specifically to genes encoding the heavy and light chains of murine antibodies). The hybridoma cells of the invention serve as a preferred source of such DNA. Once isolated, the DNA may be placed into expression 5 vectors, which are then transfected into host cells such as simian COS cells, Chinese hamster ovary (CHO) cells, or myeloma cells that do not otherwise produce immunoglobulin protein, to obtain the synthesis of monoclonal antibodies in the recombinant host cells. The DNA also may be modified, for example, by substituting the coding sequence for human heavy and light chain constant domains in place of the homologous murine sequences [U.S. Patent No. 4,816,567; Morrison *et al.*, *supra*] or by covalently joining to the immunoglobulin 10 coding sequence all or part of the coding sequence for a non-immunoglobulin polypeptide. Such a non-immunoglobulin polypeptide can be substituted for the constant domains of an antibody of the invention, or can be substituted for the variable domains of one antigen-combining site of an antibody of the invention to create a chimeric bivalent antibody.

The antibodies may be monovalent antibodies. Methods for preparing monovalent antibodies are well 15 known in the art. For example, one method involves recombinant expression of immunoglobulin light chain and modified heavy chain. The heavy chain is truncated generally at any point in the Fc region so as to prevent heavy chain crosslinking. Alternatively, the relevant cysteine residues are substituted with another amino acid residue or are deleted so as to prevent crosslinking.

In vitro methods are also suitable for preparing monovalent antibodies. Digestion of antibodies to 20 produce fragments thereof, particularly, Fab fragments, can be accomplished using routine techniques known in the art.

3. Humanized Antibodies

The anti-*patched-2* antibodies of the invention may further comprise humanized antibodies or human 25 antibodies. Humanized forms of non-human (e.g., murine) antibodies are chimeric immunoglobulins, immunoglobulin chains or fragments thereof (such as Fv, Fab, Fab', F(ab')₂ or other antigen-binding subsequences of antibodies) which contain minimal sequence derived from non-human immunoglobulin. Humanized antibodies include human immunoglobulins (recipient antibody) in which residues from a complementary determining region (CDR) of the recipient are replaced by residues from a CDR of a non-human 30 species (donor antibody) such as mouse, rat or rabbit having the desired specificity, affinity and capacity. In some instances, Fv framework residues of the human immunoglobulin are replaced by corresponding non-human residues. Humanized antibodies may also comprise residues that are found neither in the recipient antibody nor in the imported CDR or framework sequences. In general, the humanized antibody will comprise substantially all of at least one, and typically two, variable domains, in which all or substantially all of the CDR regions correspond to those of a non-human immunoglobulin and all or substantially all of the FR regions are those of a 35 human immunoglobulin consensus sequence. The humanized antibody optimally also will comprise at least a portion of an immunoglobulin constant region (Fc), typically that of a human immunoglobulin [Jones *et al.*, *Nature*, 321:522-525 (1986); Riechmann *et al.*, *Nature*, 332:323-329 (1988); and Presta, *Curr. Op. Struct. Biol.*,

2:593-596 (1992)].

Methods for humanizing non-human antibodies are well known in the art. Generally, a humanized antibody has one or more amino acid residues introduced into it from a source that is non-human. These non-human amino acid residues are often referred to as "import" residues, which are typically taken from an "import" variable domain. Humanization can be essentially performed following the method of Winter and co-workers [Jones *et al.*, *Nature*, 321:522-525 (1986); Riechmann *et al.*, *Nature*, 332:323-327 (1988); Verhoeyen *et al.*, *Science*, 239:1534-1536 (1988)], by substituting rodent CDRs or CDR sequences for the corresponding sequences of a human antibody. Accordingly, such "humanized" antibodies are chimeric antibodies (U.S. Patent No. 4,816,567), wherein substantially less than an intact human variable domain has been substituted by the corresponding sequence from a non-human species. In practice, humanized antibodies are typically human antibodies in which some CDR residues and possibly some FR residues are substituted by residues from analogous sites in rodent antibodies.

Human antibodies can also be produced using various techniques known in the art, including phage display libraries [Hoogenboom and Winter, *J. Mol. Biol.*, 227:381 (1991); Marks *et al.*, *J. Mol. Biol.*, 222:581 (1991)]. The techniques of Cole *et al.* and Boerner *et al.* are also available for the preparation of human monoclonal antibodies [(Cole *et al.*, *Monoclonal Antibodies and Cancer Therapy*, Alan R. Liss, p. 77 (1985) and Boerner *et al.*, *J. Immunol.*, 147(1):86-95 (1991)].

4. Bispecific Antibodies

Bispecific antibodies are monoclonal, preferably human or humanized, antibodies that have binding specificities for at least two different antigens. In the present case, one of the binding specificities is for the vertebrate *patched-2*, the other one is for any other antigen, and preferably for a cell-surface protein or receptor or receptor subunit.

Methods for making bispecific antibodies are known in the art. Traditionally, the recombinant production of bispecific antibodies is based on the co-expression of two immunoglobulin heavy-chain/light-chain pairs, where the two heavy chains have different specificities [Milstein and Cuello, *Nature*, 305:537-539 (1983)]. Because of the random assortment of immunoglobulin heavy and light chains, these hybridomas (quadromas) produce a potential mixture of ten different antibody molecules, of which only one has the correct bispecific structure. The purification of the correct molecule is usually accomplished by affinity chromatography steps. Similar procedures are disclosed in WO 93/08829, published 13 May 1993, and in Traunecker *et al.*, *EMBO J.*, 10:3655-3659 (1991).

Antibody variable domains with the desired binding specificities (antibody-antigen combining sites) can be fused to immunoglobulin constant domain sequences. The fusion preferably is with an immunoglobulin heavy-chain constant domain, comprising at least part of the hinge, CH2, and CH3 regions. It is preferred to have the first heavy-chain constant region (CH1) containing the site necessary for light-chain binding present in at least one of the fusions. DNAs encoding the immunoglobulin heavy-chain fusions and, if desired, the immunoglobulin light chain, are inserted into separate expression vectors, and are co-transfected into a suitable

host organism. For further details of generating bispecific antibodies see, for example, Suresh *et al.*, *Methods in Enzymology*, 121:210 (1986).

5. Heteroconjugate Antibodies

5 Heteroconjugate antibodies are also within the scope of the present invention. Heteroconjugate antibodies are composed of two covalently joined antibodies. Such antibodies have, for example, been proposed to target immune system cells to unwanted cells [U.S. Patent No. 4,676,980], and for treatment of HIV infection [WO 91/00360; WO 92/200373; EP 03089]. It is contemplated that the antibodies may be prepared *in vitro* using known methods in synthetic protein chemistry, including those involving crosslinking agents. For 10 example, immunotoxins may be constructed using a disulfide exchange reaction or by forming a thioether bond. Examples of suitable reagents for this purpose include iminothiolate and methyl-4-mercaptoputyrimidate and those disclosed, for example, in U.S. Patent No. 4,676,980.

G. Uses for anti-patched-2 Antibodies

15 The anti-patched-2 antibodies of the invention have various utilities. For example, anti-patched-2 antibodies may be used in diagnostic assays for *patched-2*, e.g., detecting its expression in specific cells, tissues, or serum. Various diagnostic assay techniques known in the art may be used, such as competitive binding assays, direct or indirect sandwich assays and immunoprecipitation assays conducted in either heterogeneous or homogeneous phases [Zola, *Monoclonal Antibodies: A Manual of Techniques*, CRC Press, Inc. (1987) pp. 147-20 158]. The antibodies used in the diagnostic assays can be labeled with a detectable moiety. The detectable moiety should be capable of producing, either directly or indirectly, a detectable signal. For example, the detectable moiety may be a radioisotope, such as ³H, ¹⁴C, ³²P, ³⁵S, or ¹²⁵I, a fluorescent or chemiluminescent compound, such as fluorescein isothiocyanate, rhodamine, or luciferin, or an enzyme, such as alkaline phosphatase, beta-galactosidase or horseradish peroxidase. Any method known in the art for conjugating the 25 antibody to the detectable moiety may be employed, including those methods described by Hunter *et al.*, *Nature*, 144: 945 (1962); David *et al.*, *Biochemistry*, 13:1014 (1974); Pain *et al.*, *J. Immunol. Meth.*, 40:219 (1981); and Nygren, *J. Histochem. and Cytochem.*, 30:407 (1982).

30 Anti-patched-2 antibodies also are useful for the affinity purification of *patched-2* from recombinant cell culture or natural sources. In this process, the antibodies against *patched-2* are immobilized on a suitable support, such a Sephadex resin or filter paper, using methods well known in the art. The immobilized antibody then is contacted with a sample containing the *patched-2* to be purified, and thereafter the support is washed with a suitable solvent that will remove substantially all the material in the sample except the *patched-2*, which is bound to the immobilized antibody. Finally, the support is washed with another suitable solvent that will release the *patched-2* from the antibody.

35 Basal cell carcinoma (BCC) is the most common human cancer. The *Hh* signaling pathway was found to activated in all BCCs. Loss of *Ptch* function is thought to lead to unregulated *Smo* activity and is responsible for about half of all BCCs. *Ptch* being a target of the *Hh* pathway itself, increases in *Ptch* mRNA levels have

been detected in BCC [Galiani, *et al.*, *Nature Genet.* 14: 78-81 (1996)] as well as in animal models of BCC. Oro *et al.*, *Science* 276: 817-821 (1997); Xie *et al.*, *Nature* 391: 90-92 (1998). Abnormal activation of *Sh* signaling, such as that which occurs in BCC, was examined to confirm whether *Ptch-2* (SEQ ID NO:2) expression was increased. As shown in Fig. 9, an *in situ* hybridization for *Ptch* (SEQ ID NO:4) and *Ptch-2* (SEQ ID NO:2) in *Smo-M2* (SEQ ID NO:16)-transgenic mice (Xie *et al.*, *supra*), while lower than *Ptch*, was still high in tumor cells. This suggests that therapeutic antibodies directed toward *Ptch-2* (SEQ ID NO:2) may be useful for the treatment of BCC.

5 Anti-*patched-2* antibodies also have utilities similar to those articulated for under the previous section "E. Uses of Patched-2". Depending on whether anti-*patched-2* antibodies will bind *patched-2* receptors so as to either inhibit *Hh* signaling (antagonist) or inhibit *patched-2* complexing with *Smo* (SEQ ID NO:17) and thereby 10 remove the normal inhibitory effect of *Smo* (SEQ ID NO:17) on *Hh* signaling (agonist) the antibody will have utilities corresponding to those articulated previously for *patched-2*.

H. *Patched-2* Antagonists

15 Several approaches may be suitably employed to create the *patched-2* antagonist and agonist compounds of the present invention. Any approach where the antagonist molecule can be targeted to the interior of the cell, which interferes or prevents wild type *patched-2* from normal operation is suitable. For example, competitive inhibitors, including mutant *patched-2* receptors which prevent wild type *patched-2* from properly binding with other proteins necessary for *Dhh* and *Hh* signaling. Additional properties of such antagonist or 20 agonist molecules are readily determinable by one of ordinary skill, such as size, charge and hydrophobicity suitable for transmembrane transport.

Where mimics or other mammalian homologues of *patched-2* are to be identified or evaluated, the cells are exposed to the test compound and compared to positive controls which are exposed only to human *patched-2*, and to negative controls which were not exposed to either the compound or the natural ligand. Where 25 antagonists or agonists of *patched-2* signal modulation are to be identified or evaluated, the cells are exposed to the compound of the invention in the presence of the natural ligand and compared to controls which are not exposed to the test compound.

30 Detection assays may be employed as a primary screen to evaluate the *Hh* signaling inhibition/enhancing activity of the antagonist/agonist compounds of the invention. The assays may also be used to assess the relative potency of a compound by testing a range of concentrations, in a range from 100 mM to 1 pM, for example, and computing the concentration at which the amount of phosphorylation or signal transduction is reduced or increased by 50% (IC₅₀) compared to controls.

35 Assays can be performed to identify compounds that affect *Hh* signaling of *patched-2* substrates. Specifically, assays can be performed to identify compounds that increase the phosphorylation activity of *patched-2* or assays can be performed to identify compounds that decrease the *Hh* signaling of *patched-2* substrates. These assays can be performed either on whole cells themselves or on cell extracts. The assays can be performed in a variety of formats, including protein-protein binding assays, biochemical screening assays,

immunoassays, cell based assays, *etc.* Such assay formats are well known in the art.

The screening assays of the present invention are amenable to high-throughput screening of chemical libraries, and are particularly suitable for identifying small molecule drug candidates.

5

(1) *Antagonist and agonist molecules*

To screen for antagonists and/or agonists of *patched-2* signaling, the assay mixture is incubated under conditions whereby, but for the presence of the candidate pharmacological agent, *patched-2* induces hedgehog signaling with a reference activity. The mixture components can be added in any order that provides for the requisite hedgehog activity. Incubation may be performed at any temperature that facilitates optimal binding, 10 typically between about 4° and 40°C, more commonly between about 15° and 40°C. Incubation periods are likewise selected for optimal binding but also minimized to facilitate rapid, high-throughput screening, and are typically between about 0.1 and 10 hours, preferably less than 5 hours, more preferably less than 2 hours. After incubation, the effect of the candidate pharmacological agent on the *patched-2* signaling is determined in any 15 convenient way. For cell-free binding-type assays, a separation step is often used to separate bound and unbound components. Separation may, for example, be effected by precipitation (*e.g.* TCA precipitation, immunoprecipitation, *etc.*), immobilization (*e.g.* on a solid substrate), followed by washing. The bound protein is conveniently detected by taking advantage of a detectable label attached to it, *e.g.* by measuring radioactive emission, optical or electron density, or by indirect detection using, *e.g.* antibody conjugates.

For example, a method of screening for suitable *patched-2* antagonists and/or agonists could involve the 20 application of *Dhh* and other hedgehog ligands. Such a screening assay could compare *in situ* hybridization in the presence and absence of the candidate antagonist and/or agonist in a *patched-2* expressing tissue as well as confirmation or absence of *patched-2* modulated cellular development. Typically these methods involve exposing an immobilized *patched-2* to a molecule suspected of binding thereto and determining the level of 25 ligand binding downstream activation of reporter constructs and/or evaluating whether or not the molecule activates (or blocks activation of) *patched-2*. In order to identify such *patched-2* binding ligands, *patched-2* can be expressed on the surface of a cell and used to screen libraries of synthetic candidate compounds or naturally-occurring compounds (*e.g.*, from endogenous sources such as serum or cells).

Suitable molecules that affect the protein-protein interaction of *patched-2* and its binding proteins 30 include fragments of the latter or small molecules, *e.g.*, peptidomimetics, which will inhibit ligand-receptor interaction. Such small molecules, which are usually less than 10 K molecular weight, are preferable as therapeutics since they are more likely to be permeable to cells, are less susceptible to degradation by various cellular mechanisms, and are not as apt to elicit an immune response as proteins. Small molecules include but are not limited to synthetic organic or inorganic compounds. Many pharmaceutical companies have extensive 35 libraries of such molecules, which can be conveniently screened by using the assays of the present invention. Non-limiting examples include proteins, peptides, glycoproteins, glycopeptides, glycolipids, polysaccharides, oligosaccharides, nucleic acids, bioorganic molecules, peptidomimetics, pharmacological agents and their metabolites, transcriptional and translation control sequences, and the like.

A preferred technique for identifying molecules which bind to *patched-2* utilizes a chimeric substrate (e.g., epitope-tagged *patched-2* or *patched-2* immunoadhesin) attached to a solid phase, such as the well of an assay plate. The binding of the candidate molecules, which are optionally labeled (e.g., radiolabeled), to the immobilized receptor can be measured. Alternatively, competition for various *Hh* pathways, especially *Dhh* 5 (SEQ ID NO:13) can be measured. In screening for antagonists and/or agonists, *patched-2* can be exposed to a *patched-2* substrate followed by the putative antagonist and/or agonist, or the *patched-2* binding protein and antagonist and/or agonist can be added simultaneously, and the ability of the antagonist and/or agonist to block *patched-2* activation can be evaluated.

10 (2) *Detection assays*

The *patched-2* polypeptides are useful in assays for identifying lead compounds for therapeutically active agents that modulate *patched-2* receptor/ligand hedgehog signaling. Specifically, lead compounds that either prevent the formation of *patched-2* signaling complexes or prevent or attenuate *patched-2* modulated hedgehog signaling (e.g., binding to *patched-2*) can be conveniently identified.

15 Various procedures known in the art may be used for identifying, evaluating or assaying the inhibition of activity of the *patched-2* proteins of the invention. As *patched-2* is believed to be a receptor for *Dhh* (SEQ ID NO:13), but also binds *Shh* (SEQ ID NO:14) and *Ihh* (SEQ ID NO:29), techniques known for use with identifying ligand/receptor modulators may also be employed with the present invention. In general, such assays involve exposing target cells in culture to the compounds and a) biochemically analyzing cell lysates to assess 20 the level and/or identity of binding; or (b) scoring phenotypic or functional changes in treated cells as compared to control cells that were not exposed to the test substance. Such screening assays are described in U.S.P. 5,602,171, U.S.P. 5,710,173, WO 96/35124 and WO 96/40276.

25 (a) *Biochemical detection techniques*

Biochemical analysis can be evaluated by a variety of techniques. One typical assay mixture which can be used with the present invention contains *patched-2* and a ligand protein with which *patched-2* is normally associated (e.g., *Dhh* (SEQ ID NO:13)) usually in an isolated, partially pure or pure form. One or both of these components may be *patched-2* to another peptide or polypeptide, which may, for example, provide or enhance protein-protein binding, improve stability under assay conditions, etc. In addition, one of the components 30 usually comprises or is coupled to a detectable label. The label may provide for direct detection by measuring radioactivity, luminescence, optical or electron density, etc., or indirect detection such as an epitope tag, an enzyme, etc. The assay mixture can additionally comprise a candidate pharmacological agent, and optionally a variety of other components, such as salts, buffers, carrier proteins, e.g. albumin, detergents, protease inhibitors, nucleic acid inhibitors, antimicrobial agents, etc., which facilitate binding, increase stability, reduce non-specific or 35 background interactions, or otherwise improve the efficiency or sensitivity of the assay.

The following detection methods may also be used in a cell-free system wherein cell lysate containing the signal transducing substrate molecule and *patched-2* is mixed with a compound of the invention. To assess

the activity of the compound, the reaction mixture may be analyzed by the SDS-PAGE technique or it may be added to substrate-specific anchoring antibody bound to a solid support, and a detection procedure as described above is performed on the separated or captured substrate to assess the presence or absence of a *patched-2* binding ligand. The results are compared to those obtained with reaction mixtures to which the compound is not 5 added. The cell-free system does not require the natural ligand or knowledge of its identity. For example, Posner *et al.* (U.S.P. 5,155,031 describes the use of insulin receptor as a substrate and rat adipocytes as target cells to demonstrate the ability of pervanadate to inhibit PTP activity. Another example, Burke *et al.*, *Biochem. Biophys. Res. Comm.* 204: 129-134 (1994) describes the use of autophosphorylated insulin receptor and recombinant PTP1B in assessing the inhibitory activity of a phosphotyrosyl mimetic.

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(i) *Whole cell detection*

A common technique involves incubating cells with *patched-2* and radiolabeled ligand, lysing the cells, separating cellular protein components of the lysate using an SDS-polyacrylamide gel (SDS-PAGE) technique, 15 in either one or two dimensions, and detecting the presence of labeled proteins by exposing X-ray film. Detection can also be effected without using radioactive labeling. In such a technique, the protein components (e.g., separated by SDS-PAGE) are transferred to a nitrocellulose membrane where the presence of *patched-2*-ligand complexes is detected using an anti-ligand antibody.

Alternatively, the anti-*patched-2* ligand antibody can be conjugated with an enzyme, such as 20 horseradish peroxidase, and detected by subsequent addition of a colorimetric substrate for the enzyme. A further alternative involves detecting the anti-*patched-2* ligand by reacting with a second antibody that recognizes anti-*patched-2* ligand, this second antibody being labeled with either a radioactive moiety or an 25 enzyme as previously described. Examples of these and similar techniques are described in Hansen *et al.*, *Electrophoresis* 14: 112-126 (1993); Campbell *et al.*, *J. Biol. Chem.* 268: 7427-7434 (1993); Donato *et al.*, *Cell Growth Diff.* 3: 258-268 (1992); Katagiri *et al.*, *J. Immunol.* 150: 585-593 (1993). Additionally, the anti-*patched-2* ligand can be detected by labeling it with a radioactive substance, followed by scanning the labeled 30 nitrocellulose to detect radioactivity or exposure of X-ray film.

Further detection methods may be developed which are preferred to those described above. Especially 35 for use in connection with high-throughput screening, it is expected that such methods would exhibit good sensitivity and specificity, extended linear range, low background signal, minimal fluctuation, compatibility with other reagents, and compatibility with automated handling systems.

The *in vivo* efficacy of the treatment of the present invention can be studied against chemically induced tumors in various rodent models. Tumor cell lines propagated in *in vitro* cell cultures can be introduced in 35 experimental rodents, e.g. mice by injection, for example by the subcutaneous route. Techniques for chemical induction of tumors in experimental animals are well known in the art.

(ii) *Kinase assays*

Because *patched-2* is a negative regulator of *Hh* signaling, which when activated by *Hh* releases the normal inhibitory effect on *Smo*, the inhibition of *patched-2* binding to *Smo* can be measured by activation of various kinase substrate associated with *Hh* signaling. When the screening methods of the present invention for 5 *patched-2* antagonists/agonists are carried out as an *ex vivo* assay, the target kinase (e.g. *fused*) can be a substantially purified polypeptide. The kinase substrate (e.g., MBP, *Gli*) is a substantially purified substrate, which in the assay is phosphorylated in a reaction with a substantially purified phosphate source that is catalyzed by the kinase. The extent of phosphorylation is determined by measuring the amount of substrate phosphorylated in the reaction. A variety of possible substrates may be used, including the kinase itself in which 10 instance the phosphorylation reaction measured in the assay is autophosphorylation. Exogenous substrates may also be used, including standard protein substrates such as myelin basic protein (MBP); yeast protein substrates; synthetic peptide substrates, and polymer substrates. Of these, MBP and other standard protein substrates may be regarded as preferred. Other substrates may be identified, however, which are superior by way of affinity for the kinase, minimal perturbation of reaction kinetics, possession of single or homogenous reaction sites, ease of 15 handling and post-reaction recover, potential for strong signal generation, and resistance or inertness to test compounds.

Measurement of the amount of substrate phosphorylated in the *ex vivo* assay of the invention may be carried out by means of immunoassay, radioassay or other well-known methods. In an immunoassay measurement, an antibody (such as a goat or mouse anti-phosphoserine/threonine antibody) may be used which 20 is specific for phosphorylated moieties formed during the reaction. Using well-known ELISA techniques, the phosphoserine/threonine antibody complex would itself be detected by a further antibody linked to a label capable of developing a measurable signal (as for example a fluorescent or radioactive label). Additionally, ELISA-type assays in microtitre plates may be used to test purified substrates. Peraldi *et al.*, *J. Biochem.* 285: 71-78 (1992); Schraag *et al.*, *Anal. Biochem.* 211: 233-239 (1993); Cleavland, *Anal. Biochem.* 190: 249-253 25 (1990); Farley, *Anal. Biochem.* 203: 151-157 (1992) and Lozaro, *Anal. Biochem.* 192: 257-261 (1991).

For example, detection schemes can measure substrate depletion during the kinase reaction. Initially, the phosphate source may be radiolabeled with an isotope such as ^{32}P or ^{33}P , and the amount of substrate phosphorylation may be measured by determining the amount of radiolabel incorporated into the substrate during the reaction. Detection may be accomplished by: (a) commercially available scintillant-containing plates 30 and beads using a beta-counter, after adsorption to a filter or a microtitre well surface, or (b) photometric means after binding to a scintillation proximity assay bead or scintillant plate. Weernink and Kijken, *J. Biochem. Biophys. Methods* 31: 49, 1996; Braunwalder *et al.*, *Anal. Biochem.* 234: 23 (1996); Kentrup *et al.*, *J. Biol. Chem.* 271: 3488 (1996) and Rusken *et al.*, *Meth. Enzymol.* 200: 98 (1991).

35 Preferably, the substrate is attached to a solid support surface by means of non-specific or, preferably, specific binding. Such attachment permits separation of the phosphorylated substrate from unincorporated, labeled phosphate source (such as adenosine triphosphate prior to signal detection. In one embodiment, the substrate may be physically immobilized prior to reaction, as through the use of NuncTM high protein binding

plate (Hanke *et al.*, *J. Biol. Chem.* 271: 695 (1996)) or Wallac ScintiStripTM plates (Braunwalder *et al.*, *Anal. Biochem.* 234: 23 (1996). Substrate may also be immobilized after reaction by capture on, for example, P81 phosphocellulose (for basic peptides), PEI/acidic molybdate resin or DEAE, or TCA precipitation onto WhatmanTM 3MM paper, Tiganis *et al.*, *Arch. Biochem. Biophys.* 325: 289 (1996); Morawetz *et al.*, *Mol. Gen. Genet.* 250: 17 (1996); Budde *et al.*, *Int J. Pharmacognosy* 33: 27 (1995) and Casnelli, *Meth. Enz.* 200: 115 (1991). Yet another possibility is the attachment of the substrate to the support surface, as by conjugation with binding partners such as glutathione and streptavidin (in the case of GST and biotin), respectively) which have been attached to the support, or via antibodies specific for the tags which are likewise attached to the support.

10 Further detection methods may be developed which are preferred to those described above. Especially for use in connection with high-throughput screening, it is expected that such methods would exhibit good sensitivity and specificity, extended linear range, low background signal, minimal fluctuation, compatibility with other reagents, and compatibility with automated handling systems.

15 The *in vivo* efficacy of the treatment of the present invention can be studied against chemically induced tumors in various rodent models. Tumor cell lines propagated in *in vitro* cell cultures can be introduced in experimental rodents, *e.g.* mice by injection, for example by the subcutaneous route. Techniques for chemical induction of tumors in experimental animals are well known in the art.

| (b) *Biological detection techniques:*

20 The ability of the antagonist/agonist compounds of the invention to modulate the activity of *patched-2*, which itself modulates hedgehog signaling, may also be measured by scoring for morphological or functional changes associated with ligand binding. Any qualitative or quantitative technique known in the art may be applied for observing and measuring cellular processes which comes under the control of *patched-2*. The activity of the compounds of the invention can also be assessed in animals using experimental models of disorders caused by or related to dysfunctional *hedgehog* signaling. For example, ineffective *Dhh* hedgehog 25 signaling in mice leads to viable but sterile mice. Additionally, proper *Shh* signaling is critical to murine embryonic development at the notochord and floor plate, neural tube, distal limb structures, spinal column and ribs. Improper *Shh* signaling, is also correlative with cyclopia. Any of these phenotypic properties could be evaluated and quantified in a screening assay for *patched-2* antagonists and/or agonist. Disease states associated 30 with overexpression of hedgehog is associated with basal cell carcinoma while inactive *Shh* signaling leads to improper neural development.

| The data obtained from these cell culture assays and animal studies can be used in formulating a range of dosages for use in humans. The dosage of the compounds of the invention should lie within a range of circulating concentrations with little or no toxicity. The dosage may vary within this range depending on the dosage form employed and the route of administration.

35

| (2) *Antisense oligonucleotides*

| Another preferred class of antagonists involves the use of gene therapy techniques, include-including

the administration of antisense oligonucleotides. Applicable gene therapy techniques include single or multiple administrations of therapeutically effective DNA or mRNA. Antisense RNAs and DNAs can be used as therapeutic agents for blocking the expression of certain genes *in vivo*. Reference short antisense oligonucleotides can be imported into cells where they act as inhibitors, despite their low intracellular concentrations caused by restricted uptake by the cell membrane, Zamecnik *et al.*, *Proc. Natl. Acad. Sci. USA* 83: 4143-4146 (1986). The anti-sense oligonucleotides can be modified to enhance their uptake, *e.g.*, by substituting their negatively charged phosphodiester groups by uncharged groups.

There are a variety of techniques known for introducing nucleic acids into viable cells. The techniques vary depending upon whether the nucleic acid is transferred into cultured cells *in vitro*, *ex vivo*, or *in vivo* in the cells of the intended host. Techniques suitable for the transfer of nucleic acid into mammalian cells *in vitro* include the use of liposomes, electroporation, microinjection, cell fusion, DEAE-dextran, the calcium phosphate precipitation method, *etc.* The currently preferred *in vivo* gene transfer techniques include transfection with viral (typically retroviral) vectors and viral coat protein-liposome mediated transfection, Dzau *et al.*, *Trends Biotech.* 11: 205-210 (1993). In some situations it is desirable to provide the nucleic acid source with an agent that targets the cells, such as an antibody specific for a cell surface membrane protein associated with endocytosis may be used for targeting and/or to facilitate uptake, *e.g.*, capsid proteins or fragments thereof tropic for a particular cell type, antibodies for proteins which undergo internalization in cycling, and proteins that target intracellular localization and enhance intracellular half-life. The technique of receptor-mediated endocytosis is described, for example, by Wu *et al.*, *J. Biol. Chem.* 262: 4429-4432 (1987); Wagner *et al.*, *Proc. Natl. Acad. Sci. USA* 87: 3410-3414 (1990). For a review of known gene targeting and gene therapy protocols, see Anderson *et al.*, *Science* 256: 808-813 (1992).

In one embodiment of the invention, *patched-2* expression may be reduced by providing *patched-2*-expressing cells with an amount of *patched-2* antisense RNA or DNA effective to reduce expression of the *patched-2* protein.

25

I. Diagnostic Uses

Another use of the compounds of the invention (*e.g.*, *patched-2*, *patched-2* variant and anti-*patched-2* antibodies) described herein is to help diagnose whether a disorder is driven, to some extent, by *patched-2* or hedgehog signaling. For example, basal cell carcinoma cells are associated with active *hedgehog* signaling, spermatocyte formation is associated with *Dhh* signaling, and defective *Ptch* and *Ptch-2* suppression may be associated with testicular carcinomas.

A diagnostic assay to determine whether a particular disorder is driven by *Ptch-2* modulated *hedgehog* signaling, can be carried out using the following steps: (1) culturing test cells or tissues; (2) administering a compound which can prevent *Ptch-2* (SEQ ID NO:2)-binding with *Smo* (SEQ ID NO:17), thereby activating the *Hh* signaling pathway; and (3) measuring the amount of *Hh* signaling. The steps can be carried out using

standard techniques in light of the present disclosure. For example, standard techniques can be used to isolate cells or tissues and culturing *or in vivo*.

Compounds of varying degree of selectivity are useful for diagnosing the role of *patched-2*. For example, compounds which inhibit *patched-2* in addition to another form of kinase can be used as an initial test compound to determine if one of several signaling ligands drive the disorder. The selective compounds can then be used to further eliminate the possible role of the other ligands in driving the disorder. Test compounds should be more potent in inhibiting ligand-*patched-2* binding activity than in exerting a cytotoxic effect (*e.g.*, an IC₅₀/LD₅₀ of greater than one). The IC₅₀ and LD₅₀ can be measured by standard techniques, such as an MTT assay, or by measuring the amount of LDH released. The degree of IC₅₀/LD₅₀ of a compound should be taken into account in evaluating the diagnostic assay. For example, the larger the IC₅₀/LD₅₀ ratio the more relative the information. Appropriate controls take into account the possible cytotoxic effect of a compound of a compound, such as treating cells not associated with a cell proliferative disorder (*e.g.*, control cells) with a test compound, can also be used as part of the diagnostic assay. The diagnostic methods of the invention involve the screening for agents that modulate the effects of *patched-2* upon *hedgehog* signaling. Exemplary detection techniques include radioactive labeling and immunoprecipitating (U.S.P. 5,385,915).

J. Pharmaceutical Compositions and Dosages

Therapeutic formulations of the compositions of the invention are prepared for storage as lyophilized formulations or aqueous solutions by mixing the *patched-2* molecule, agonist and/or antagonist having the desired degree of purity with optional "pharmaceutically-acceptable" or "physiologically-acceptable" carriers, excipients or stabilizers typically employed in the art (all of which are termed "excipients"). For example, buffering agents, stabilizing agents, preservatives, isotonifiers, non-ionic detergents, antioxidants and other miscellaneous additives. (See Remington's Pharmaceutical Sciences, 16th Ed., A. Osol, Ed. (1980)). Such additives must be nontoxic to the recipients at the dosages and concentrations employed.

Buffering agents help to maintain the pH in the range which approximates physiological conditions. They are preferably present at concentration ranging from about 2mM to about 50 mM. Suitable buffering agents for use with the present invention include both organic and inorganic acids and salts thereof such as citrate buffers (*e.g.*, monosodium citrate-disodium citrate mixture, citric acid-trisodium citrate mixture, citric acid-monosodium citrate mixture, *etc.*), succinate buffers (*e.g.*, succinic acid-monosodium succinate mixture, succinic acid-sodium hydroxide mixture, succinic acid-disodium succinate mixture, *etc.*), tartrate buffers (*e.g.*, tartaric acid-sodium tartrate mixture, tartaric acid-potassium tartrate mixture, tartaric acid-sodium hydroxide mixture, *etc.*), fumarate buffers (*e.g.*, fumaric acid-monosodium fumarate mixture, *etc.*), fumarate buffers (*e.g.*, fumaric acid-monosodium fumarate mixture, fumaric acid-disodium fumarate mixture, monosodium fumarate-disodium fumarate mixture, *etc.*), gluconate buffers (*e.g.*, gluconic acid-sodium glyconate mixture, gluconic acid-sodium hydroxide mixture, gluconic acid-potassium glyconate mixture, *etc.*), oxalate buffer (*e.g.*, oxalic acid-sodium oxalate mixture, oxalic acid-sodium hydroxide mixture, oxalic acid-potassium oxalate mixture, *etc.*), lactate buffers (*e.g.*, lactic acid-sodium lactate mixture, lactic acid-sodium hydroxide mixture, lactic acid-

potassium lactate mixture, *etc.*) and acetate buffers (*e.g.*, acetic acid-sodium acetate mixture, acetic acid-sodium hydroxide mixture, *etc.*). Additionally, there may be mentioned phosphate buffers, histidine buffers and trimethylamine salts such as Tris.

5 Preservatives are added to retard microbial growth, and are added in amounts ranging from 0.2% - 1% (w/v). Suitable preservatives for use with the present invention include phenol, benzyl alcohol, *meta*-cresol, methyl paraben, propyl paraben, octadecyldimethylbenzyl ammonium chloride, benzalconium halides (*e.g.*, chloride, bromide, iodide), hexamethonium chloride, alkyl parabens such as methyl or propyl paraben, catechol, resorcinol, cyclohexanol, and 3-pentanol.

10 Isotonifiers sometimes known as "stabilizers" are present to ensure isotonicity of liquid compositions of the present invention and include polyhydric sugar alcohols, preferably trihydric or higher sugar alcohols, such as glycerin, erythritol, arabitol, xylitol, sorbitol and mannitol. Polyhydric alcohols can be present in an amount between 0.1% to 25% by weight, preferably 1% to 5% taking into account the relative amounts of the other ingredients.

15 Stabilizers refer to a broad category of excipients which can range in function from a bulking agent to an additive which solubilizes the therapeutic agent or helps to prevent denaturation or adherence to the container wall. Typical stabilizers can be polyhydric sugar alcohols (enumerated above); amino acids such as arginine, lysine, glycine, glutamine, asparagine, histidine, alanine, ornithine, L-leucine, 2-phenylalanine, glutamic acid, threonine, *etc.*, organic sugars or sugar alcohols, such as lactose, trehalose, stachyose, mannitol, sorbitol, xylitol, ribitol, myoinisitol, galactitol, glycerol and the like, including cyclitols such as inositol; polyethylene glycol; 20 amino acid polymers; sulfur containing reducing agents, such as urea, glutathione, thiocitic acid, sodium thioglycolate, thioglycerol, α -monothioglycerol and sodium thio sulfate; low molecular weight polypeptides (*i.e.* < 10 residues); proteins such as human serum albumin, bovine serum albumin, gelatin or immunoglobulins; hydrophilic polymers, such as polyvinylpyrrolidone monosaccharides, such as xylose, mannose, fructose, glucose; disaccharides such as lactose, maltose, sucrose and trisaccacharides such as raffinose; polysaccharides 25 such as dextran. Stabilizers can be present in the range from 0.1 to 10,000 weights per part of weight active protein.

Non-ionic surfactants or detergents (also known as "wetting agents") are present to help solubilize the therapeutic agent as well as to protect the therapeutic protein against agitation-induced aggregation, which also permits the formulation to be exposed to shear surface stressed without causing denaturation of the protein. 30 Suitable non-ionic surfactants include polysorbates (20, 80, *etc.*), polyoxamers (184, 188 *etc.*), Pluronic[®] polyols, polyoxyethylene sorbitan monoethers (Tween[®]-20, Tween[®]-80, *etc.*). Non-ionic surfactants are present in a range of about 0.05 mg/ml to about 1.0 mg/ml, preferably about 0.07 mg/ml to about 0.2 mg/ml.

Additional miscellaneous excipients include bulking agents, (*e.g.* starch), chelating agents (*e.g.* EDTA), antioxidants (*e.g.*, ascorbic acid, methionine, vitamin E), and cosolvents.

35 The formulation herein may also contain more than one active compound as necessary for the particular indication being treated, preferably those with complementary activities that do not adversely affect each other.

For example, it may be desirable to further provide an immunosuppressive agent. Such molecules are suitably present in combination in amounts that are effective for the purpose intended.

The active ingredients may also be entrapped in microcapsule prepared, for example, by coacervation techniques or by interfacial polymerization, for example, hydroxymethylcellulose or gelatin-microcapsule and poly-(methylmethacrylate) microcapsule, respectively, in colloidal drug delivery systems (for example, liposomes, albumin microspheres, microemulsions, nano-particles and nanocapsules) or in macroemulsions. Such techniques are disclosed in *Remington's Pharmaceutical Sciences*, 16th edition, A. Osal, Ed. (1980).

The formulations to be used for in vivo administration must be sterile. This is readily accomplished, for example, by filtration through sterile filtration membranes.

Sustained-release preparations may be prepared. Suitable examples of sustained-release preparations include semi-permeable matrices of solid hydrophobic polymers containing the compounds of the invention, which matrices are in the form of shaped articles, *e.g.*, films, or microcapsules. Examples of sustained-release matrices include polyesters, hydrogels (for example, poly(2-hydroxyethyl-methacrylate), or poly(vinylalcohol)), polylactides (U.S. Pat. No.3,773,919), copolymers of L-glutamic acid and ethyl-L-glutamate, non-degradable ethylene-vinyl acetate, degradable lactic acid-glycolic acid copolymers such as the LUPRON DEPOT™ (injectable microspheres composed of lactic acid-glycolic acid copolymer and leuprolide acetate), and poly-D-(β)-3-hydroxybutyric acid. While polymers such as ethylene-vinyl acetate and lactic acid-glycolic acid enable release of molecules for over 100 days, certain hydrogels release proteins for shorter time periods. When encapsulated compounds of the invention remain in the body for a long time, they may denature or aggregate as a result of exposure to moisture at 37°C, resulting in a loss of biological activity and possible changes in immunogenicity. Rational strategies can be devised for stabilization depending on the mechanism involved. For example, if the aggregation mechanism is discovered to be intermolecular S-S bond formation through thio-disulfide interchange, stabilization may be achieved by modifying sulfhydryl residues, lyophilizing from acidic solutions, controlling moisture content, using appropriate additives, and developing specific polymer matrix compositions.

The amount of therapeutic polypeptide, antibody or fragment thereof which will be effective in the treatment of a particular disorder or condition will depend on the nature of the disorder or condition, and can be determined by standard clinical techniques. Where possible, it is desirable to determine the dose-response curve and the pharmaceutical compositions of the invention first *in vitro*, and then in useful animal model systems prior to testing in humans. However, based on common knowledge of the art, a pharmaceutical composition effective in modulating *Dhh* and *Hh* signaling may provide a local *patched-2* protein concentration of between about 10 and 1000 ng/ml, preferably between 100 and 800 ng/ml and most preferably between about 200 ng/ml and 600 ng/ml of *Ptch-2* (SEQ ID NO:2).

In a preferred embodiment, an aqueous solution of therapeutic polypeptide, antibody or fragment thereof is administered by subcutaneous injection. Each dose may range from about 0.5 μ g to about 50 μ g per kilogram of body weight, or more preferably, from about 3 μ g to about 30 μ g per kilogram body weight.

The dosing schedule for subcutaneous administration may vary from once a week to daily depending on a number of clinical factors, including the type of disease, severity of disease, and the subject's sensitivity to the therapeutic agent.

5 *Patched-2* polypeptide may comprise an amino acid sequence or subsequence thereof as indicated in Fig. 1 (SEQ ID NO:2), active amino acid sequence derived therefrom, or functionally equivalent sequence as this subsequence is believed to comprise the functional portion of the *patched-2* polypeptide.

10 If the subject manifests undesired side effects such as temperature elevation, cold or flu-like symptoms, fatigue, etc., it may be desirable to administer a lower dose at more frequent intervals. One or more additional drugs may be administered in combination with *patched-2* to alleviate such undesired side effects, for example, an anti-pyretic, anti-inflammatory or analgesic agent.

The following examples are offered for illustrative purposes only, and are not intended to limit the scope of the present invention in any way.

15 All patent and literature references cited in the present specification are hereby incorporated by reference in their entirety.

EXAMPLES

20 Commercially available reagents referred to in the examples were used according to manufacturer's instructions unless otherwise indicated. The source of those cells identified in the following examples, and throughout the specification, by ATCC accession numbers is the American Type Culture Collection, Rockville, Maryland.

EXAMPLE 1

25 **Introduction:**

At the cell surface, *Hh* function appears to be mediated by a multicomponent receptor complex involving *Ptch* (SEQ ID NO:4) and *Smo* (SEQ ID NO:17), two multi-transmembrane proteins initially identified as segment polarity genes in *Drosophila* and later characterized in vertebrates. Nakano, Y. *et al.*, *Nature* 341: 508-513 (1989); Goodrich *et al.*, *Gene Dev.* 10: 301-312 (1996); Marigo *et al.*, *Develop.* 122: 1225-1233 (1996); van 30 den Heuvel *et al.*, *Nature* 382: 547-551 (1996); Alcedo *et al.*, *Cell* 86: 221-232 (1996); Stone *et al.* *Nature* 384: 129-34 (1996). Both genetic and biochemical evidence support the existence of a receptor complex where *Ptch* (SEQ ID NO:4) is the ligand binding subunit, and where *Smo* (SEQ ID NO:17), a G-protein coupled receptor like molecule, is the signaling component. Stone *et al.*, *Nature* 384: 129-134 (1996), Marigo *et al.*, *Nature* 384: 176-79 (1996), Chen *et al.*, *Cell* 87: 553-63 (1996). Upon binding of *Hh* to *Ptch* (SEQ ID NO:4), the normal 35 inhibitory effect of *Ptch* (SEQ ID NO:4) on *Smo* (SEQ ID NO:17) is relieved, allowing *Smo* (SEQ ID NO:17) to transduce the *Hh* signal across the plasma membrane.

Results:

It remains to be established if the *patched-Smoothened* receptor complex mediates the action of all 3 mammalian Hhs or if specific components exist. Recently, a second murine *patched* gene, *Ptch-2* (SEQ ID NO:7) was recently isolated [Motoyama *et al.*, *Nature Genet.* 18: 104-106 (1998)] but its function as a *Hh* receptor has not been established. In order to characterize *Ptch-2* (SEQ ID NO:2) and compare it to *Ptch* (SEQ ID NO:4) with respect to the biological function of the various *Hh* family members, we have screened EST databases with the *Ptch* (SEQ ID NO:4) protein and identified 2 EST candidates for a novel human *patched* gene. A full length cDNA encoding human *Ptch-2* (SEQ ID NO:2) was cloned from a testis library. The initiation ATG defines a 3612 nucleotide open reading frame encoding a 1204 amino acid long protein with a predicted molecular weight of approximately 131 kDa. The overall identity between human *Ptch* (SEQ ID NO:4) (SEQ ID NO:7) and *Ptch-2* (SEQ ID NO:2) is 54% (Fig. 1), while the identity between human *Ptch-2* (SEQ ID NO:2) and the recently described mouse *Ptch-2* (SEQ ID NO:7) (SEQ ID NO:8) is 90%. (Fig. 8). The most obvious structural difference between the two human *Ptch* proteins is a truncated C-terminal cytoplasmic domain in *Ptch-2* (SEQ ID NO:2). In addition, only one of the two glycosylation sites present in *Ptch* (SEQ ID NO:4) is conserved in *Ptch-2* (SEQ ID NO:2).

To determine if *Ptch-2* (SEQ ID NO:2) is a *Hh* receptor and if the two *patched* molecules are capable of discriminating between the various *Hh* ligands through specific binding, Applicants transfected human 293 embryonic kidney cells with *Ptch* (SEQ ID NO:4) or *Ptch-2* (SEQ ID NO:2)-expression constructs and analyzed the cells for binding of *Shh*, *Dhh* and *Ihh* (SEQ ID NOS:14, 13, and 29, respectively). As shown on Figure 7A, binding of ^{125}I -*Shh* can be competed with an excess of *Shh*, *Dhh* or *Ihh* (SEQ ID NOS:14, 13, and 29, respectively). Scatchard analysis of the displacement curves indicates that all *Hhs* have similar affinity for *Ptch* (SEQ ID NO:4) (*Shh* (SEQ ID NO:14), 1.0nM; *Dhh* (SEQ ID NO:13), 2.6nM; *Ihh* (SEQ ID NO:29), 1.0nM) and *Ptch-2* (SEQ ID NO:2) (*Shh* (SEQ ID NO:14), 1.8nM; *Dhh* (SEQ ID NO:13), 0.6nM; *Ihh* (SEQ ID NO:29), 0.4nM) indicating that both *Ptch* (SEQ ID NO:4) and *Ptch-2* (SEQ ID NO:2) can serve as physiological receptors for the 3 mammalian *Hh* proteins.

Applicants next determined whether, like *Ptch* (SEQ ID NO:4), *Ptch-2* (SEQ ID NO:2) forms a physical complex with *Smo* (SEQ ID NO:17). Expression constructs for Flag-tagged *Ptch* (SEQ ID NO:4) or *Ptch-2* (SEQ ID NO:2) were transiently co-transfected in 293 cells with Myc-tagged *Smo* (SEQ ID NO:17). As described previously [Stone *et al.*, *Nature* 384: 129-34 (1996)], in cells expressing *Ptch* (SEQ ID NO:4) and *Smo* (SEQ ID NO:17), *Ptch* (SEQ ID NO:4) can be immunoprecipitated with antibodies against the epitope-tagged *Smo* (SEQ ID NO:15) (Fig. 7B). Similarly, *Ptch-2* (SEQ ID NO:2) can be immunoprecipitated with antibodies against the epitope-tagged *Smo* (SEQ ID NO:15) when the two proteins are co-expressed in 293 cells. Together, these results suggest a model where *Ptch-2* (SEQ ID NO:2) forms a multicomponent *Hh* receptor complex with *Smo* (SEQ ID NO:17) similar to the one described for *patched* (Stone *et al.*, *supra*). Interestingly, these results also demonstrate that the long C-terminal tail which is missing in *Ptch-2* (SEQ ID NO:2) is not required for the interaction with *Smo* (SEQ ID NO:17) as was already suggested by the analysis of truncated *patched* (Stone *et al.*, *supra*). However, it remains possible that the absence of a C-terminal domain

affects the capacity of *Ptch-2* (SEQ ID NO:2) to block signaling by *Smo* or leads to difference in signaling by *patched* compared to *patched-2*.

To further investigate whether *patched-2* could mediate the action of a specific Hh molecule based on its expression profile, Applicants have compared the expression pattern of *Ptch* (SEQ ID NO:4) and *Ptch-2* (SEQ ID NO:2). First, Northern blot analysis using a probe specific for *Ptch-2* (SEQ ID NO:1) revealed high levels of *patched-2* mRNA in the testis (Fig. 4). By this method, *Ptch-2* (SEQ ID NO:1) expression was not detected in any other tissue analyzed including embryonic tissues (data not shown). This profile is very different from the one observed for *Ptch* (SEQ ID NO:18) which was not found in testis by Northern blot but in a large number of adult and embryonic tissues [Goodrich *et al.*, *Genes Dev.* 10: 301-312 (1996)]. More detailed analysis of the expression pattern of *Ptch* (SEQ ID NO:18) and *Ptch-2* (SEQ ID NO:1) was performed by *in situ* hybridization with particular attention to testis. As previously described (Motoyama *et al.*, *supra*), low levels of *Ptch-2* (SEQ ID NO:1) expression were detected in epithelial cells of the developing tooth and skin (data not shown). High levels of *Ptch-2* (SEQ ID NO:2) are expressed inside the seminiferous tubule, on the primary and secondary spermatocytes (Fig. 6B,6E) while only low levels of *Ptch* (SEQ ID NO:4) can be detected on the Leydig cells located in the interstitium of the seminiferous tubules (Fig. 6A). The primary and secondary spermatocytes are in close contact with the supporting Sertoli cells, the source of *Dhh* (SEQ ID NO:13) in the testis [Bitgood *et al.*, *Curr. Biol.* 6: 298-304 (1996)]. To determine which one of the 2 receptors is the most relevant mediator of *Dhh* (SEQ ID NO:13)-activity in the testis, we have analyzed the expression profile of FuRK (SEQ ID NO:10), a Fused Related Kinase that was shown to be a component of the *Hh* signaling pathway (Zhang *et al.*, submitted; copending U.S.S.N. Serial Number 09/031,563, filed 26 Feb. 1998). Consistent with the idea that *Ptch-2* (SEQ ID NO:2) is the target of *Dhh* in the testis, we found that FuRK (SEQ ID NO:10) is expressed only in germ cells where it colocalizes with *Ptch-2* (SEQ ID NO:2) (Figure 4c,f). *Dhh* (SEQ ID NO:13) is required for proper differentiation of germ cells since male *Dhh*-deficient mice are sterile due to lack of mature sperm (Bitgood *et al.*, *supra*). Our data suggest that *Dhh* (SEQ ID NO:13) acts directly on germ cells through *Ptch-2* (SEQ ID NO:2) while the function of *Ptch* (SEQ ID NO:4) expressed at low levels on testosterone producing Leydig cells is unclear.

Discussion:

Loss of heterozygosity (LOH) for *patched* was reported to occur with high frequency in familial as well as sporadic basal cell carcinoma [Johnson *et al.*, *Science* 272: 1668-71 (1996); Hahn *et al.*, *Cell* 85: 841-51 (1996); Gailani *et al.*, *Nature Genetics* 14: 78-81; Xie *et al.*, *Cancer Res.* 57: 2369-72 (1997)], suggesting that it functions as a tumor suppressor. According to the receptor model described above, loss of *patched* function may result in aberrant signaling by *Smo* (SEQ ID NO:17), leading to hyperproliferation of the skin basal cell layer. If, as suggested above, *patched-2* mediates the function of *Dhh*, loss of *Ptch-2* (SEQ ID NO:2) may lead to tumor formation in tissues where *Smo* (SEQ ID NO:17)-activity is controlled by *patched-2*. The gene encoding *patched-2* was mapped by fluorescence *in situ* hybridization and by PCR using a radiation hybrid

panel to human chromosome 1p33-34 (data not shown). Interestingly, recent analysis of recurrent chromosomal abnormalities in testicular tumors, including seminomas, revealed a deletion of the region 1p32-36 [Summersgill *et al.*, *B. J. Cancer* 77: 305-313 (1998)]. Loss of this region encompassing the *patched-2* locus was consistent in 36% of the germ cell tumor cases. These data raise the possibility that, like *patched* in basal cell carcinoma and medulloblastoma, *patched-2* may be a tumor suppressor in *Dhh* (SEQ ID NO:13)-target cells such as spermatocytes, further implicating *Hh* signaling in cancer.

In summary, our data demonstrate that both *patched* and *patched-2* are genuine *Hh* receptors and that they are both capable of forming a complex with *Smo* (SEQ ID NO:17). Although binding data indicate that *patched* and *patched-2* do not discriminate between the various *Hh* ligands through affinity differences, the distinct tissue distribution of these 2 receptors suggests that *in vivo*, *patched* may be the primary receptor for *Shh* whereas *Ptch-2* will mediate mainly *Dhh* signaling. The function of *patched* expression in Leydig cells in the absence of some of the *Hh* signaling components remain to be explained. Similarly, it will be of interest to determine if *patched-2* plays a role when expressed in *Shh* expressing cells present in the developing tooth and skin Motoyama *et al.*, *Nature Genet.* 18: 104-106 (1998). Finally, the existence of *patched-2* raises the question of whether additional *patched* receptors exist, in particular one that mediates the function of *Ihh* (SEQ ID NO:29).

Material and Methods:

20 1. *Isolation of human patched-2 cDNA clones*

An expressed sequence tag (EST) DNA database (LIFESEQ®, Incyte Pharmaceuticals, Palo Alto, CA) was searched for a human homologue of the *Drosophila* segment polarity gene *patched-2*. Two ESTs (Incyte #905531 and 1326258) (Fig. 2) (Figures 2A & 2B) (SEQ ID NOS: 3 and 45, respectively) were identified as a potential candidates. In order to identify human cDNA libraries containing human *patched-2* clones, human cDNA libraries in pRK5 were first screened by PCR using the following primers:

5'-905531(A): 5'-AGGCAGGGGATCACAGCA-3' (SEQ ID NO:19) (SEQ ID NO:11)

3'-905531(A): 5'-ATACCAAAGAGTTCCACT-3' (SEQ ID NO:20) (SEQ ID NO:12)

A fetal lung library was selected and enriched for *patched-2* cDNA clones by extension of single stranded DNA from plasmid libraries grown in dut/ung host using the 3'-905531(A) primer in a reaction containing 10μl of 10x PCR Buffer (Klentaq®), 1μl dNTP (200 μM), 1 μl library DNA (200 ng), 0.5 μl primer, 86.5 μl H₂O and 1 μl of Klentaq® (Clontech) added after a hot start. The reaction was denatured for 1 min. at 95°C, annealed for 1 min. at 60°C then extended for 20 min. at 72°C. DNA was extracted with phenol/CHCl₃, ethanol precipitated, then transformed by electroporation into DH10B (Gibco/BRL) host bacteria. Colonies from each transformation were replica plated on nylon membranes and screened with an overlapping oligo probe derived from the EST sequence (#905531) of the following sequence:

5'-*Ptch2* probe: 5'-CTGCGGCGCTGCTCCTGCTGGCCGCTGCATCCTGCTGGTGTGC-3' (SEQ ID NO:21) (SEQ ID NO:13)

3'-*Ptch2* probe: 5'-AGAGCACAGACGAGGAAAGTGCACACCAGCAGGATGCAGACGGCC-3' (SEQ ID NO:22) (SEQ ID NO:14)

5 The oligo probe was labeled with [γ -³²P]-ATP and T4 polynucleotide kinase. Filters were hybridized overnight at 42°C in 50% formamide, 5xSSC, 10xDenhardt's, 0.05M sodium phosphate (pH 6.5), 0.1% sodium pyrophosphate, 50 μ g/ml of sonicated salmon sperm DNA. The filters were then rinsed in 2x SSC and washed in 0.1x SSC, 0.1% SDS then exposed to Kodak[®] X Ray films.

10 Using this procedure, a partial clone was isolated from the fetal brain library (clone 3A – Fig. 10) (SEQ ID NO:8) (SEQ ID NO:9). In order to isolate the missing 5'-sequence, a testis library (see northern blot analysis, infra) was screened. The primer set used to amplify a 204 bp probe from clone 3A to probe the testis library was:

RACE 5: 5'-ACTCCTGACTTGTAGCAGATT-3' (SEQ ID NO:23) (SEQ ID NO:15) and

RACE 6: 5'-AGGCTGCATACACCTCTCAGA-3'. (SEQ ID NO:24) (SEQ ID NO:16)

15 The amplified probe was purified by excision from an agarose gel and labeled with a random primer labeling kit (Boehringer Mannheim). Several clones were isolated, including one (clone 16.1 – Fig. 11 (SEQ ID NO:9)) containing a potential initiation methionine. A full length cDNA encoding *patched-2* was reconstructed by assembling several of these clones. The full length cDNA encoding human *Ptch-2* (Fig. 1) (SEQ ID NO:1) has a 3612 nucleotide long open reading frame encoding a 1204 amino acid protein with a 144 kDa predicted 20 molecular weight. Alignment with human *Ptch* (SEQ ID NO:4) reveals a 53% identity between the 2 molecules at the amino acid level (Fig. 3). All 12 transmembrane domains are conserved. The most significant difference is a shorter C-terminal intracellular domain in *Ptch-2* (SEQ ID NO:2) compared to *Ptch* (SEQ ID NO:4).

2. *Northern blot analysis:*

25

In order to determine the best tissue source for isolation of the complete full length *Ptch-2* cDNA as well as to determine its expression profile, we probed human multiple tissue northern blots (Clontech) with a 752 bp fragment amplified from the 3' untranslated region of *Ptch-2* (SEQ ID NO:2) using the following primers:

TM2: TM2 5-GCTTAGGCCGAGGAGAT-3' (SEQ ID NO:25) (SEQ ID NO:17)

30

UTR2: 5'-AACTCACAACTTCTCTCCA-3'. (SEQ ID NO:26) (SEQ ID NO:18)

35

The resulting fragment was gel purified and labeled by random priming. The blots were hybridized in ExpressHyb[®] hybridization solution (Clontech) in the presence of 1 x 10⁶ cpm/ml ³²P-labeled probe at 42°C overnight. The blots were washed in 2x SSC at room temperature for 10 minutes and washed in 0.1 x SSC/0.1 % SDS at 42°C for 30 minutes then exposed to x-ray film overnight. Fig. 4 shows that *Ptch-2* message is expressed at high levels in only the testis.

3. *Chromosomal localization:*

The primers TM2 (SEQ ID NO:25) (SEQ ID NO:17) and UTR2 (SEQ ID NO:26) (SEQ ID NO:18) described above were used to screen the Genome Systems (St. Louis, MO) BAC library. Two individual BAC clones were obtained from this library and chromosomal localization of both of the clones by FISH indicated that *Ptch-2* (SEQ ID NO:2) maps to human chromosome 1p33-34 (FIG 5). Loss of heterozyosity (LOH) for *patched* was reported to occur with high frequency in basal cell carcinoma. Loss of *patched* function is thought to lead to constitutive signaling by *Smoothened* (*Smo* (SEQ ID NO:17)), resulting in hyperproliferation of the basal layer of the dermis. A similar mechanism may lead to the formation of germ cell tumors. This model proposes that the first step in the progression of a germ cell tumor is an initial loss of DNA by a germ cell precursor, leading to a neoplastic germ cell which then forms a seminoma [De Jong *et al.*, *Cancer Genet. Cytogenet.* 48: 143-167 (1990)]. From the invasive seminoma, all other forms of germ cell tumor types develop. Approximately 80% of all germ cell tumors correlate with an isochromosome 12p (i12p) and is found at a higher frequency in non-seminomas than seminomas [Rodriguez *et al.*, *Cancer Res.* 52: 2285-2291 (1992)]. However, analysis of recurrent chromosomal abnormalities in testicular tumors including seminomas revealed a deletion of the region 1p32-36. Loss of this region was consistent in 36% of the germ cell tumor cases of in a recent study Summersgill *et al.*, *B. J. Cancer* 57: 305-313 (1998)]. A similar deletion of chromosome 1p32-36 has been reported at a frequency of 28% in oligodendrogiomas Bello, *et al.*, *Int. J. Cancer* 57: 172-175 (1994). While expression of *patched-2* (SEQ ID NO:2) in the brain was not examined here in detail, *patched-2* (SEQ ID NO:2) is thought to be the *Dhh* receptor (see below) and expression of *Dhh* by murine Schwann cells was previously reported [Bitgood *et al.*, *Develop. Biol.* 172: 126-138 (1995)]. Since *patched-2* (SEQ ID NO:2) localizes to chromosome 1p33-34 it is possible that *patched-2* regulates *Smo* (SEQ ID NO:17) signaling in *Dhh* target cells and that loss of *patched-2* function leads to abnormal *Smo* (SEQ ID NO:17) signaling in these cells and subsequent tumor formation.

25 4. *In situ hybridization:*

Mouse testis sections were cut at 16 μ m, and processed for in situ hybridization by the method described in Phillips *et al.*, *Science* 250: 290-294 (1990). 33 P-UTP labeled RNA probes were generated as described in Melton *et al.*, *Nucleic Acids Res.* 12: 7035-7052 (1984). Sense and antisense probes were synthesized from the 3' non coding region of the mouse *Ptch* or *Ptch-2* and from a mouse FuRK cDNA fragment corresponding to the region encoding amino acid 317-486 of the human sequence using T3 and T7, respectively.

Ptch:

503 (Anti-sense) 5'GGATTCTAATACGACTCACTATAGGGCCAATGGCCTAACCGACTGC3' (SEQ ID NO:27) (SEQ ID NO:19)

503 (Sense)

5'CTATGAAATTAACCCTCACTAAAGGGACCCACGGCCTCTCCTCAC3' (SEQ ID NO:28) (SEQ ID
NO:20)

Ptch2:

504 (Anti-sense)

5 5'GGATTCTAATACGACTCACTATAGGGCCCCTAAACTCCGCTGCTCCAC3' (SEQ ID NO:12) (SEQ ID
NO:21)

504 (Sense)

5'CTATGAAATTAACCCTCACTAAAGGGAGCTCCGTGAGTCCCTATGTG3' (SEQ ID NO:11) (SEQ ID
NO:22)

10

FuRK sense and antisense were synthesized from a mouse *fused* DNA fragment using T3 and T7, respectively, corresponding to the region encoding amino acid residues 317-486 of the human sequence (Zhang *et al.*, submitted, 1998; copending U.S.S.N. 09/031,563, filed 26 Feb. 1998).

15

Figure 6 illustrates that, although both *Ptch* (SEQ ID NO:4) and *Ptch-2* (SEQ ID NO:2) are expressed in testis, their expression pattern does not overlap. *Ptch* (SEQ ID NO:4) is expressed in the Leydig cells of the interstitium while *Ptch-2* (SEQ ID NO:2) is expressed in the primary and secondary spermatocytes.

20

The expression of *Ptch-2* (SEQ ID NO:2) specifically in the developing spermatogonia suggest that *Ptch-2* (SEQ ID NO:2) is the immediate target of *Dhh* (SEQ ID NO:13). *Dhh* (SEQ ID NO:13) is expressed by Sertoli cells and mice deficient in *Dhh* (SEQ ID NO:13) are sterile because of a defect in sperm production [Bitgood *et al.*, *Curr. Biol.* 6: 298-304 (1996)]. Although this effect on germ cells was thought to be indirect and mediated by *Ptch* present on Leydig cells, our data suggest that *Dhh* directly acts on germ cells through *Ptch-2*. This is further demonstrated by the localization of FuRK (SEQ ID NO:10), an intracellular kinase homologous to *Drosophila Fused* and involved in transducing the Hedgehog (*Hh*) signal. As shown in Figure 6, FuRK (SEQ ID NO:10) is colocalizes with *Ptch-2* (SEQ ID NO:2) in germ cells and not with *Ptch* (SEQ ID NO:4) in Leydig cells, suggesting that *Ptch-2* and not *Ptch* will be able to transduce the *Dhh* signal. These results suggest that *Ptch-2* is a *Dhh* receptor.

25

30 *Ptch-2* mRNA levels in *Smo-M2* (SEQ ID NO:16) transgenic mice [A *Smo* mutation which results in autonomous phenotypes similar to BCC, Xie *et al.*, *Nature* 391: 90-92 (1998)] can be increased upon abnormal activation of the *Hh* signaling pathway. As indicated in Fig. 9, *Ptch-2* levels were high in tumor cells (although lower than *Ptch* levels). This suggests that antibodies directed toward *Ptch-2* may be useful in the treatments of BCC.

5. *Immunoprecipitation with Smo:*

The binding of *Ptch-2* to *Smo* (SEQ ID NO:17) was assessed by cotransfection using a transient transfection system of a myc-epitope tagged *Smo* (SEQ ID NO:15) and a FLAG-epitope tagged *Ptch* or *Ptch2* expression construct in 293 cells using standard techniques (Gorman, C., *DNA Cloning: A Practical Approach*, 5 Clover, DM ed., Vol. 11, pp. 143-190, IRL Press, Washington, D.C.). 36 hours after transfection, the cells were lysed in 1% NP-40 and immunoprecipitated overnight with the 9E10 anti-myc antibody or with the M2 anti-FLAG antibody (IBI-Kodak) followed by protein A Sepharose, and then separated on a denatured 6% polyacrylamide gel. Proteins were detected by transfer to nitrocellulose and probing with antibodies to Flag or Myc epitopes, using the ECL detection system (Amersham). Figure 7B indicates that both *Ptch* (SEQ ID NO:4) and *Ptch-2* (SEQ ID NO:2) are expressed at the same level (IP Flag, Blot Flag) and that like *Ptch* (SEQ ID NO:4), *Ptch-2* (SEQ ID NO:2) forms a physical complex with *Smo* (SEQ ID NO:17). These results suggest that like *patched*, *patched-2* controls *Hh* signaling through its interaction with *Smo* (SEQ ID NO:17).

10 6. *Hh Binding:*

15 To determine whether *Ptch-2* is able to bind to the various hedgehog ligands, 293 cells were transfected with *Ptch* (SEQ ID NO:4) or *Ptch-2* (SEQ ID NO:2) using standard procedures. Cells were incubated with 100 pM 125 I-*Shh* (19kD amino terminal fragment of murine *Shh* (SEQ ID NO:14)) in the presence or absence of excess unlabeled *Shh* (SEQ ID NO:14) or *Dhh* (SEQ ID NO:13) for 2h at room 20 temperature. After equilibrium was reached, the ligand bound cells were centrifuged through a continuous sucrose gradient to separate unincorporated and then counted in a scintillation counter. Figure 7A shows that both *Dhh* (SEQ ID NO:13) and *Shh* (SEQ ID NO:14) bind to *Ptch* (SEQ ID NO:4) and *Ptch-2* (SEQ ID NO:2). Varying concentrations of cold competitor indicate that the 2 ligands have similar affinity for *Ptch* (SEQ ID NO:4) and *Ptch-2* (SEQ ID NO:2).

25

Example 2

Expression of *patched-2* in *E. coli*

The DNA sequence encoding human *patched-2* is initially amplified using selected PCR primers. The primers should contain restriction enzyme sites that correspond to the restriction enzyme sites on the selected expression vector. A variety of expression vectors may be employed. An example of a suitable vector is 30 pBR322 (derived from *E. coli*; see Bolivar *et al.*, *Gene*, 2:95 (1977)) which contains genes for ampicillin and tetracycline resistance. The vector is digested with restriction enzyme and dephosphorylated. The PCR amplified sequences are then ligated into the vector. The vector will preferably include sequences that encode for an antibiotic resistance gene, a trp promoter, a polyhis leader (including the first six STII codons, polyhis sequence, and enterokinase cleavage site), the vertebrate *patched-2* coding region, lambda transcriptional 35 terminator, and an argU gene.

The ligation mixture is then used to transform a selected *E. coli* strain using the methods described in Sambrook *et al.*, *supra*. Transformants are identified by their ability to grow on LB plates and antibiotic

resistant colonies are then selected. Plasmid DNA can be isolated and confirmed by restriction analysis and DNA sequencing.

Selected clones can be grown overnight in liquid culture medium such as LB broth supplemented with antibiotics. The overnight culture may subsequently be used to inoculate a larger scale culture. The cells are then grown to a desired optical density, during which the expression promoter is turned on.

After culturing the cells for several more hours, the cells can be harvested by centrifugation. The cell pellet obtained by the centrifugation can be solubilized using various agents known in the art, and the solubilized vertebrate *patched-2* protein can then be purified using a metal chelating column under conditions that allow tight binding of the protein.

10

Example 3

Expression of patched-2 in mammalian cells

The vector, pRK5 (see EP 307,247, published March 15, 1989), is employed as the expression vector. Optionally, the vertebrate *patched-2* DNA is ligated into pRK5 with selected restriction enzymes to allow 15 insertion of the vertebrate *patched-2* DNA using ligation methods such as described in Sambrook *et al.*, *supra*. The resulting vector is called pRK5-patched-2.

In one embodiment, the selected host cells may be 293 cells. Human 293 cells (ATCC CCL 1573) are grown to confluence in tissue culture plates in medium such as DMEM supplemented with fetal calf serum and optionally, nutrient components and/or antibiotics. About 10 µg pRK5-patched-2 DNA is mixed with about 1 µg 20 DNA encoding the VA RNA gene [Thimmappaya *et al.*, *Cell*, 31:543 (1982)] and dissolved in 500 µl of 1 mM Tris-HCl, 0.1 mM EDTA, 0.227 M CaCl₂. To this mixture is added, dropwise, 500 µl of 50 mM HEPES (pH 7.35), 280 mM NaCl, 1.5 mM NaPO₄, and a precipitate is allowed to form for 10 minutes at 25°C. The precipitate is suspended and added to the 293 cells and allowed to settle for about four hours at 37°C. The culture medium is aspirated off and 2 ml of 20% glycerol in PBS is added for 30 seconds. The 293 cells are then 25 washed with serum free medium, fresh medium is added and the cells are incubated for about 5 days.

Approximately 24 hours after the transfections, the culture medium is removed and replaced with culture medium (alone) or culture medium containing 200 µCi/ml ³⁵S-cysteine and 200 µCi/ml ³⁵S-methionine. After a 12 hour incubation, the conditioned medium is collected, concentrated on a spin filter, and loaded onto a 30 15% SDS gel. The processed gel may be dried and exposed to film for a selected period of time to reveal the presence of vertebrate *patched-2* polypeptide. The cultures containing transfected cells may undergo further incubation (in serum free medium) and the medium is tested in selected bioassays.

In an alternative technique, vertebrate *patched-2* may be introduced into 293 cells transiently using the dextran sulfate method described by Somparyrac *et al.*, *Proc. Natl. Acad. Sci.*, 78:7575 (1981). 293 cells are grown to maximal density in a spinner flask and 700 µg pRK5-patched-2 DNA is added. The cells are first 35 concentrated from the spinner flask by centrifugation and washed with PBS. The DNA-dextran precipitate is incubated on the cell pellet for four hours. The cells are treated with 20% glycerol for 90 seconds, washed with

tissue culture medium, and re-introduced into the spinner flask containing tissue culture medium, 5 μ g/ml bovine insulin and 0.1 μ g/ml bovine transferrin. After about four days, the conditioned media is centrifuged and filtered to remove cells and debris. The sample containing expressed vertebrate *patched-2* can then be concentrated and purified by any selected method, such as dialysis and/or column chromatography.

5 In another embodiment, vertebrate *patched-2* can be expressed in CHO cells. The pSVi-*patched-2* can be transfected into CHO cells using known reagents such as CaPO₄ or DEAE-dextran. As described above, the cell cultures can be incubated, and the medium replaced with culture medium (alone) or medium containing a radiolabel such as ³⁵S-methionine. After determining the presence of vertebrate *patched-2* polypeptide, the culture medium may be replaced with serum free medium. Preferably, the cultures are incubated for about 6
10 days, and then the conditioned medium is harvested. The medium containing the expressed vertebrate *patched-2* can then be concentrated and purified by any selected method.

Epitope-tagged vertebrate *patched-2* may also be expressed in host CHO cells. The vertebrate *patched-2* may be subcloned out of the pRK5 vector. The subclone insert can undergo PCR to fuse in frame with a selected epitope tag such as a poly-his tag into an expression vector. The poly-his tagged vertebrate *patched-2* insert can then be subcloned into a SV40 driven vector containing a selection marker such as DHFR for selection
15 of stable clones. Finally, the CHO cells can be transfected (as described above) with the SV40 driven vector. Labeling may be performed, as described above, to verify expression. The culture medium containing the expressed poly-His tagged vertebrate *patched-2* can then be concentrated and purified by any selected method, such as by Ni²⁺-chelate affinity chromatography.

20

Example 4

Expression of vertebrate *patched-2* in Yeast

The following method describes recombinant expression of vertebrate *patched-2* in yeast.

First, yeast expression vectors are constructed for intracellular production or secretion of vertebrate
25 *patched-2* from the ADH2/GAPDH promoter. DNA encoding vertebrate *patched-2*, a selected signal peptide and the promoter is inserted into suitable restriction enzyme sites in the selected plasmid to direct intracellular expression of vertebrate *patched-2*. For secretion, DNA encoding vertebrate *patched-2* can be cloned into the selected plasmid, together with DNA encoding the ADH2/GAPDH promoter, the yeast alpha-factor secretory signal/leader sequence, and linker sequences (if needed) for expression of vertebrate *patched-2*.

30 Yeast cells, such as yeast strain AB110, can then be transformed with the expression plasmids described above and cultured in selected fermentation media. The transformed yeast supernatants can be analyzed by precipitation with 10% trichloroacetic acid and separation by SDS-PAGE, followed by staining of the gels with Coomassie Blue stain.

35 Recombinant vertebrate *patched-2* can subsequently be isolated and purified by removing the yeast cells from the fermentation medium by centrifugation and then concentrating the medium using selected cartridge filters. The concentrate containing vertebrate *patched-2* may further be purified using selected column chromatography resins.

Example 5

Expression of vertebrate *patched-2* in Baculovirus-Infected Insect Cells

The following method describes recombinant expression of vertebrate *patched-2* in Baculovirus-infected insect cells.

5 The vertebrate *patched-2* is *patched-2* upstream of an epitope tag contained within a baculovirus expression vector. Such epitope tags include poly-his tags and immunoglobulin tags (like Fc regions of IgG). A variety of plasmids may be employed, including plasmids derived from commercially available plasmids such as pVL1393 (Novagen). Briefly, the vertebrate *patched-2* or the desired portion of the vertebrate *patched-2* (such as the sequence encoding the extracellular domain of a transmembrane protein) is amplified by PCR with 10 primers complementary to the 5' and 3' regions. The 5' primer may incorporate flanking (selected) restriction enzyme sites. The product is then digested with those selected restriction enzymes and subcloned into the expression vector.

15 Recombinant baculovirus is generated by co-transfected the above plasmid and BaculoGold™ virus DNA (Pharmingen) into *Spodoptera frugiperda* ("Sf9") cells (ATCC CRL 1711) using lipofectin (commercially available from GIBCO-BRL). After 4 - 5 days of incubation at 28°C, the released viruses are harvested and used for further amplifications. Viral infection and protein expression is performed as described by O'Reilley *et al.*, Baculovirus expression vectors: A laboratory Manual, Oxford: Oxford University Press (1994).

20 Expressed poly-his tagged vertebrate *patched-2* can then be purified, for example, by Ni²⁺-chelate affinity chromatography as follows. Extracts are prepared from recombinant virus-infected Sf9 cells as described by Rupert *et al.*, *Nature*, 362:175-179 (1993). Briefly, Sf9 cells are washed, resuspended in sonication buffer (25 mL Hepes, pH 7.9; 12.5 mM MgCl₂; 0.1 mM EDTA; 10% Glycerol; 0.1% NP-40; 0.4 M KCl), and sonicated twice for 20 seconds on ice. The sonicates are cleared by centrifugation, and the supernatant is diluted 50-fold in loading buffer (50 mM phosphate, 300 mM NaCl, 10% Glycerol, pH 7.8) and filtered through a 0.45 µm filter. A Ni²⁺-NTA agarose column (commercially available from Qiagen) is prepared with a bed volume of 25 5 mL, washed with 25 mL of water and equilibrated with 25 mL of loading buffer. The filtered cell extract is loaded onto the column at 0.5 mL per minute. The column is washed to baseline A₂₈₀ with loading buffer, at which point fraction collection is started. Next, the column is washed with a secondary wash buffer (50 mM phosphate; 300 mM NaCl, 10% Glycerol, pH 6.0), which elutes nonspecifically bound protein. After reaching A₂₈₀ baseline again, the column is developed with a 0 to 500 mM Imidazole gradient in the secondary wash buffer. One mL fractions are collected and analyzed by SDS-PAGE and silver staining or western blot with Ni²⁺-NTA-conjugated to alkaline phosphatase (Qiagen). Fractions containing the eluted His₁₀-tagged vertebrate *patched-2* are pooled and dialyzed against loading buffer.

30 Alternatively, purification of the IgG tagged (or Fc tagged) vertebrate *patched-2* can be performed using known chromatography techniques, including for instance, Protein A or protein G column chromatography.

Example 6

Preparation of Antibodies that Bind Vertebrate *patched-2*

This example illustrates preparation of monoclonal antibodies, which can specifically bind vertebrate
5 *patched-2*.

Techniques for producing the monoclonal antibodies are known in the art and are described, for instance, in Goding, *supra*. Immunogens that may be employed include purified vertebrate *patched-2*, fusion proteins containing vertebrate *patched-2*, and cells expressing recombinant vertebrate *patched-2* on the cell surface. Selection of the immunogen can be made by the skilled artisan without undue experimentation.

10 Mice, such as Balb/c, are immunized with the vertebrate *patched-2* immunogen (E.g., extracellular portions or cells expressed *Ptch-2*) emulsified in complete Freund's adjuvant and injected subcutaneously or intraperitoneally in an amount from 1-100 micrograms. Alternatively, the immunogen is emulsified in MPL-TDM adjuvant (Ribi Immunochemical Research, Hamilton, MT) and injected into the animal's hind foot pads. The immunized mice are then boosted 10 to 12 days later with additional immunogen emulsified in the selected
15 adjuvant. Thereafter, for several weeks, the mice may also be boosted with additional immunization injections. Serum samples may be periodically obtained from the mice by retro-orbital bleeding for testing in ELISA assays to detect vertebrate *patched-2* antibodies.

20 After a suitable antibody titer has been detected, the animals "positive" for antibodies can be injected with a final intravenous injection of vertebrate *patched-2*. Three to four days later, the mice are sacrificed and the spleen cells are harvested. The spleen cells are then patched-2 (using 35% polyethylene glycol) to a selected murine myeloma cell line such as P3X63AgU.1, available from ATCC, No. CRL 1597. The fusions generate hybridoma cells which can then be plated in 96 well tissue culture plates containing HAT (hypoxanthine, aminopterin, and thymidine) medium to inhibit proliferation of non-patched-2 cells, myeloma hybrids, and spleen cell hybrids.

25 The hybridoma cells will be screened in an ELISA for reactivity against vertebrate *patched-2*. Determination of "positive" hybridoma cells secreting the desired monoclonal antibodies against vertebrate *patched-2* is within the skill in the art.

30 The positive hybridoma cells can be injected intraperitoneally into syngeneic Balb/c mice to produce ascites containing the anti- vertebrate *patched-2* monoclonal antibodies. Alternatively, the hybridoma cells can be grown in tissue culture flasks or roller bottles. Purification of the monoclonal antibodies produced in the ascites can be accomplished using ammonium sulfate precipitation, followed by gel exclusion chromatography. Alternatively, affinity chromatography based upon binding of antibody to protein A or protein G can be employed.

Example 7

Gli Luciferase Assay

5 The following assay may be used to measure the activation of the transcription factor *GLI*, the mammalian homologue of the *Drosophila cubitus interruptus* (Ci). It has been shown that *GLI* is a transcription factor activated upon *SHh* stimulation of cells.

Nine (9) copies of a *GLI* binding site present in the *HNF3β* enhancer, (Sasaki *et al.*, *Development* 124: 1313-1322 (1997)), are introduced in front of a thymidine kinase minimal promoter driving the luciferase reporter gene in the pGL3 plasmid (Promega). The sequence of the *GLI* binding sequence is: 10 TCGACAAGCAGGAAACACCCAAGTAGAAGCTC (p9XGliLuc) ~~(SEQ ID NO:31)~~ (SEQ ID NO:23), while the negative control sequence is: TCGACAAGCAGGGAAGTGGGAAGT ~~AGTAGAAGCTC~~ (p9XmGliLuc) ~~(SEQ ID NO:32)~~ (SEQ ID NO:24). These constructs are cotransfected with the full length *Ptch-2* and *Smo* in C3H10T1/2 cells grown in F12, DMEM (50:50), 10% FCS heat inactivated. The day before transfection 1 x 10⁵ cells per well was inoculated in 6 well plates, in 2 ml of media. The following day, 1 µg of each construct is 15 cotransfected in duplicate with 0.025 mg ptkRenilla luciferase plasmid using lipofectamine (Gibco-BRL) in 100 µl OptiMem (with GlutaMAX) as per manufacturer's instructions for 3 hours at 37°C. Serum (20%, 1 ml) is then added to each well and the cells were incubated for 3 more hours at 37°C. Cells are then washed twice with PBS, then incubated for 48 hours at 37°C in 2 ml of media. Each well is then washed with PBS, and the cells 20 lysed in 0.5 ml Passive Lysis Buffer (Promega) for 15 min. at room temperature on a shaker. The lysate is transferred in eppendorf tubes on ice, spun in a refrigerated centrifuge for 30 seconds and the supernatant saved on ice. For each measure, 20 µl of cell lysate is added to 100 µl of LARII (luciferase assay reagent, Promega) in a polypropylene tube and the luciferase light activity measured. The reaction is stopped by the addition of Stop and Glow buffer (Promega), mixed by pipetting up and down 3 to 5 times and *Renilla* luciferase lights activity is measured on the luminometer.

* * * *

Deposit of Material

The following materials have been deposited with the American Type Culture Collection, 12301 Parklawn Drive, Rockville, MD, USA (ATCC):

5	<u>Designation:</u>	<u>ATCC Dep. No.</u>	<u>Deposit Date</u>
	pRK7.hptc2.Flag-1405	209778	4/14/98

10 This deposit was made under the provisions of the Budapest Treaty on the International Recognition of the Deposit of Microorganisms for the Purpose of Patent Procedure and the Regulations thereunder (Budapest Treaty). This assures maintenance of a viable culture of the deposit for 30 years from the date of deposit. The deposit will be made available by ATCC under the terms of the Budapest Treaty, and subject to an agreement between Genentech, Inc. and ATCC, which assures permanent and unrestricted availability of the progeny of the culture of the deposit to the public upon issuance of the pertinent U.S. patent or upon laying open to the public 15 of any U.S. or foreign patent application, whichever comes first, and assures availability of the progeny to one determined by the U.S. Commissioner of Patents and Trademarks to be entitled thereto according to 35 USC §122 and the Commissioner's rules pursuant thereto (including 37 CFR §1.14 with particular reference to 886 OG 638).

20 The assignee of the present application has agreed that if a culture of the materials on deposit should die or be lost or destroyed when cultivated under suitable conditions, the materials will be promptly replaced on notification with another of the same. Availability of the deposited material is not to be construed as a license to practice the invention in contravention of the rights granted under the authority of any government in accordance with its patent laws.

25 The foregoing written specification is considered to be sufficient to enable one skilled in the art to practice the invention. The present invention is not to be limited in scope by the construct deposited, since the deposited embodiment is intended as a single illustration of certain aspects of the invention and any constructs that are functionally equivalent are within the scope of this invention. The deposit of material herein does not constitute an admission that the written description herein contained is inadequate to enable the practice of any aspect of the invention, including the best mode thereof, nor is it to be construed as limiting the scope of the 30 claims to the specific illustrations that it represents. Indeed, various modifications of the invention in addition to those shown and described herein will become apparent to those skilled in the art from the foregoing description and fall within the scope of the appended claims.

Sequence Listing

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Substitute Specification

Patent Docket No. P1405R1C1

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80	Gly Pro Phe Ala Ser Leu Glu Gly Phe Arg Glu Leu Leu Asp Lys	235	240
	230	235	
85	Ala Gln Val Gly Gln Ala Tyr Val Gly Arg Pro Cys Leu Asp Pro	250	255
	245	250	
90	Asp Asp Pro His Cys Pro Pro Ser Ala Pro Asn Arg His Ser Arg	265	270
	260	265	
95	Gln Ala Pro Asn Val Ala Gln Glu Leu Ser Gly Gly Cys His Gly	280	285
	275	280	

~~Phe Ser His Lys Phe Met His Trp Cln Glu Glu Leu Leu Leu Gly~~
290 ~~295~~ 300

5 ~~Gly Thr Ala Arg Asp Leu Cln Gly Cln Leu Leu Arg Ala Glu Ala~~
305 ~~310~~ 315

~~Leu Cln Ser Thr Phe Leu Leu Met Ser Pro Arg Cln Leu Tyr Glu~~
320 ~~325~~ 330

10 ~~His Phe Arg Gly Asp Tyr Cln Thr His Asp Ile Gly Trp Ser Glu~~
335 ~~340~~ 345

Substitute Specification

Patent Docket No. P1405R1C1

Glu Cln Ala Ser Met Val Leu Cln Ala Trp Gln Arg Arg Phe Val
350 355 360

Gln Leu Ala Gln Glu Ala Leu Pro Ala Asn Ala Ser Cln Gln Ile
365 370 375

His Ala Phe Ser Ser Thr Thr Leu Asp Asp Ile Leu Arg Ala Phe
380 385 390

Ser Glu Val Ser Thr Thr Arg Val Val Cln Gly Tyr Leu Leu Met
395 400 405

Leu Ala Tyr Ala Cys Val Thr Met Leu Arg Trp Asp Cys Ala Cln
410 415 420

Ser Gln Gly Ala Val Gly Leu Ala Gly Val Leu Leu Val Ala Leu
425 430 435

Ala Val Ala Ser Gly Leu Gly Leu Cys Ala Leu Leu Gly Ile Thr
440 445 450

Phe Asn Ala Ala Thr Thr Gln Val Leu Pro Phe Leu Ala Leu Cln
455 460 465

Ile Cln Val Asp Asp Ile Phe Leu Leu Ala His Ala Phe Thr Lys
470 475 480

Ala Pro Pro Asp Thr Pro Leu Pro Cln Arg Met Cln Cln Cys Leu
485 490 495

Arg Ser Thr Gly Thr Ser Val Ala Leu Thr Ser Val Asn Asn Met
500 505 510

Val Ala Phe Phe Met Ala Ala Leu Val Pro Ile Pro Ala Leu Arg
515 520 525

Ala Phe Ser Leu Cln Ala Ala Ile Val Val Cln Cys Asn Phe Ala
530 535 540

Ala Val Met Leu Val Phe Pro Ala Ile Leu Ser Leu Asp Leu Arg
545 550 555

Arg Arg His Arg Cln Arg Leu Asp Val Leu Cln Cys Cys Phe Ser Ser
560 565 570

Pro Cys Ser Ala Cln Val Ile Cln Met Leu Pro Cln Glu Leu Cln
575 580 585

Asp Arg Ala Val Pro Val Gly Ile Ala His Leu Thr Ala Thr Val
590 595 600

Gln Ala Phe Thr His Cys Glu Ala Ser Ser Gln His Val Val Thr
5 605 610 615

Ile Leu Pro Pro Cln Ala His Leu Leu Ser Pro Ala Ser Asp Pro
620 625 630

Leu Gly Ser Glu Leu Tyr Ser Pro Gly Gly Ser Thr Arg Asp Leu
10 635 640 645

Leu Ser Gln Glu Glu Gly Thr Gly Pro Gln Ala Ala Cys Arg Pro
15 650 655 660

Leu Leu Cys Ala His Trp Thr Leu Ala His Phe Ala Arg Tyr Gln
665 670 675

Phe Ala Pro Leu Leu Leu Gln Thr Arg Ala Lys Ala Leu Val Leu
20 680 685 690

Leu Phe Phe Gly Ala Leu Leu Gly Leu Ser Leu Tyr Gly Ala Thr
695 700 705

Leu Val Gln Asp Gly Leu Ala Leu Thr Asp Val Val Pro Arg Gly
25 710 715 720

Thr Lys Glu His Ala Phe Leu Ser Ala Gln Leu Arg Tyr Phe Ser
725 730 735

Leu Tyr Glu Val Ala Leu Val Thr Gln Gly Gly Phe Asp Tyr Ala
30 740 745 750

His Ser Gln Arg Ala Leu Phe Asp Leu His Gln Arg Phe Ser Ser
35 755 760 765

Leu Lys Ala Val Leu Pro Pro Ala Thr Gln Ala Pro Arg Thr
770 775 780

Trp Leu His Tyr Tyr Arg Ser Trp Leu Gln Gly Ile Gln Ala Ala
40 785 790 795

Phe Asp Gln Asp Trp Ala Ser Gly Arg Ile Thr Cys His Ser Tyr
800 805 810

Arg Asn Gly Ser Glu Asp Gly Ala Leu Ala Tyr Lys Leu Leu Ile
45 815 820 825

1 Gln Thr Gly Asn Ala Gln Glu Pro Leu Asp Phe Ser Gln Leu Thr
2 830 835 840

5 Thr Arg Lys Leu Val Asp Lys Glu Gly Leu Ile Pro Pro Glu Leu
6 845 850 855

10 Phe Tyr Met Gly Leu Thr Val Trp Val Ser Ser Asp Pro Leu Gly
11 860 865 870

15 Leu Ala Ala Ser Gln Ala Asn Phe Tyr Pro Pro Pro Pro Glu Trp
16 875 880 885

20 Leu His Asp Lys Tyr Asp Thr Thr Gly Glu Asn Leu Arg Ile Pro
21 890 895 900

25 Ala Ala Gln Pro Leu Glu Phe Ala Gln Phe Pro Phe Leu Leu His
26 905 910 915

30 Gly Leu Gln Lys Thr Ala Asp Phe Val Glu Ala Ile Glu Gly Ala
31 920 925 930

35 Arg Ala Ala Cys Thr Glu Ala Gly Gln Ala Gly Val His Ala Tyr
36 935 940 945

40 Pro Ser Gly Ser Pro Phe Leu Phe Trp Glu Gln Tyr Leu Gly Leu
41 950 955 960

45 Arg Arg Cys Phe Leu Leu Ala Val Cys Ile Leu Leu Val Cys Thr
46 965 970 975

50 Phe Leu Val Cys Ala Leu Leu Leu Ser Pro Trp Thr Ala Gly
51 980 985 990

55 Leu Ile Val Leu Val Leu Ala Met Met Thr Val Glu Leu Phe Gly
56 995 1000 1005

60 Ile Met Gly Phe Leu Gly Ile Lys Leu Ser Ala Ile Pro Val Val
61 1010 1015 1020

65 Ile Leu Val Ala Ser Ile Gly Ile Gly Val Glu Phe Thr Val His
66 1025 1030 1035

70 Val Ala Leu Gly Phe Leu Thr Ser His Gly Ser Arg Asn Leu Arg
71 1040 1045 1050

75 Ala Ala Ser Ala Leu Glu Gln Thr Phe Ala Pro Val Thr Asp Gly
76 1055 1060 1065

Ala Val Ser Thr Leu Leu Gly Leu Leu Met Leu Ala Gly Ser Asn
1070 1075 1080

5 Phe Asp Phe Ile Ile Arg Tyr Phe Phe Val Val Leu Thr Val Leu
1085 1090 1095

Thr Leu Leu Gly Leu Leu His Gly Leu Leu Leu Leu Pro Val Leu
1100 1105 1110

10 Leu Ser Ile Leu Gly Pro Pro Pro Cln Val Val Cln Val Tyr Lys
1115 1120 1125

15 Glu Ser Pro Cln Thr Leu Asn Ser Ala Ala Pro Cln Arg Gly Gly
1130 1135 1140

Leu Arg Trp Asp Arg Pro Pro Thr Leu Pro Cln Ser Phe Ala Arg
1145 1150 1155

20 Val Thr Thr Ser Met Thr Val Ala Leu His Pro Pro Pro Leu Pro
1160 1165 1170

Gly Ala Tyr Val His Pro Ala Ser Glu Glu Pro Thr
1175 1180 1182

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<211> 4004

<212> DNA

<213> Homo sapiens

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gaaatggcc tccacatcaa ggcageeeete aetgeeagta aagtcataat 150

atcaatctat gggaaatgtt gggatttgaa caaaatctgt tacaaggatcg 200

gaggttttttt tattggaaat ggaatgttgg agggatgtat tgagaatgtg 250

40 ttttgttgcg tgatcttca ccccttcgac tggttttggg agggagccaa 300

actttttttt ggtttttttt acatggggatc aatggggatc tttttttttt 350

45 gagttttttt tggatgttgc tttttttttt atttatgtttt tttttttttt 400

aattttttttt gggatgttgc gttttttttt cttttttttt tttttttttt 450

50 gttttttttt tttttttttt tttttttttt tttttttttt tttttttttt 500

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Leu Glu Asp Asp Gly Lys Leu Pro Glu Asp Gln Val Gln Ala Ile
95 100 105

5 Ala Ala Gln Leu Val Ser Ala Leu Tyr Tyr Leu His Ser His Arg
110 115 120

Ile Leu His Arg Asp Met Lys Pro Gln Asn Ile Leu Leu Ala Lys
125 130 135

10 Gly Gly Gly Ile Lys Leu Cys Asp Phe Gly Phe Ala Arg Ala Met
140 145 150

Ser Thr Asn Thr Met Val Leu Thr Ser Ile Lys Gly Thr Pro Leu
155 160 165

Tyr Met Ser Pro Glu Leu Val Glu Glu Arg Pro Tyr Asp His Thr
170 175 180

20 Ala Asp Leu Trp Ser Val Gly Cys Ile Leu Tyr Glu Leu Ala Val
185 190 195

Gly Thr Pro Pro Phe Tyr Ala Thr Ser Ile Phe Gln Leu Val Ser
200 205 210

25 Leu Ile Leu Lys Asp Pro Val Arg Trp Pro Ser Thr Ile Ser Pro
215 220 225

Cys Phe Lys Asn Phe Leu Gln Gly Leu Leu Thr Lys Asp Pro Arg
30 230 235 240

Gln Arg Leu Ser Trp Pro Asp Leu Leu Tyr His Pro Phe Ile Ala
245 250 255

35 Gly His Val Thr Ile Ile Thr Glu Pro Ala Gly Pro Asp Leu Gly
260 265 270

Thr Pro Phe Thr Ser Arg Leu Pro Pro Glu Leu Cln Val Leu Lys
275 280 285

Asp Glu Cln Ala His Arg Leu Ala Pro Lys Gly Asn Cln Ser Arg
290 295 300

Ile Leu Thr Cln Ala Tyr Lys Arg Met Ala Glu Glu Ala Met Cln
305 310 315

Lys Lys His Cln Asn Thr Gly Pro Ala Leu Glu Cln Glu Asp Lys
320 325 330

Thr Ser Lys Val Ala Pro Gly Thr Ala Pro Leu Pro Arg Leu Gly
335 340 345

Ala Thr Pro Cln Glu Ser Ser Leu Leu Ala Gly Ile Leu Ala Ser
350 355 360

Glu Leu Lys Ser Ser Trp Ala Lys Ser Gly Thr Gly Glu Val Pro
365 370 375

Ser Ala Pro Arg Glu Asn Arg Thr Thr Pro Asp Cys Glu Arg Ala
380 385 390

Phe Pro Glu Glu Arg Pro Glu Val Leu Gly Cln Arg Ser Thr Asp
395 400 405

Val Val Asp Leu Glu Asn Glu Glu Pro Asp Ser Asp Asn Glu Trp
410 415 420

Cln His Leu Leu Glu Thr Thr Glu Pro Val Pro Ile Cln Leu Lys
425 430 435

Ala Pro Leu Thr Leu Leu Cys Asn Pro Asp Phe Cys Cln Arg Ile
440 445 450

Cln Ser Cln Leu His Glu Ala Gly Gly Cln Ile Leu Lys Gly Ile
455 460 465

Leu Glu Gly Ala Ser His Ile Leu Pro Ala Phe Arg Val Leu Ser
470 475 480

Ser Leu Leu Ser Ser Cys Ser Asp Ser Val Ala Leu Tyr Ser Phe
485 490 495

Cys Arg Glu Ala Gly Leu Pro Gly Leu Leu Leu Ser Leu Leu Arg
500 505 510

His Ser Gln Clu Ser Asn Ser Leu Gln Gln Ser Trp Tyr Gly
515 520 525

5 Thr Phe Leu Gln Asp Leu Met Ala Val Ile Gln Ala Tyr Phe Ala
530 535 540

Cys Thr Phe Asn Leu Glu Arg Ser Gln Thr Ser Asp Ser Leu Gln
545 550 555

10 Val Phe Gln Glu Ala Ala Asn Leu Phe Leu Asp Leu Leu Gly Lys
560 565 570

Leu Leu Ala Gln Pro Asp Asp Ser Glu Gln Thr Leu Arg Arg Asp
575 580 585

15 Ser Leu Met Cys Phe Thr Val Leu Cys Glu Ala Met Asp Gly Asn
590 595 600

Ser Arg Ala Ile Ser Lys Ala Phe Tyr Ser Ser Leu Leu Thr Thr
20 605 610 615

Gln Gln Val Val Leu Asp Gly Leu Leu His Gly Leu Thr Val Pro
620 625 630

25 Gln Leu Pro Val His Thr Pro Gln Gly Ala Pro Gln Val Ser Gln
635 640 645

Pro Leu Arg Glu Gln Ser Glu Asp Ile Pro Gly Ala Ile Ser Ser
30 650 655 660

Ala Leu Ala Ala Ile Cys Thr Ala Pro Val Gly Leu Pro Asp Cys
665 670 675

35 Trp Asp Ala Lys Glu Gln Val Cys Trp His Leu Ala Asn Gln Leu
680 685 690

Thr Glu Asp Ser Ser Gln Leu Arg Pro Ser Leu Ile Ser Gly Leu
40 695 700 705

Gln His Pro Ile Leu Cys Leu His Leu Leu Lys Val Leu Tyr Ser
710 715 720

45 Cys Cys Leu Val Ser Glu Gly Leu Cys Arg Leu Leu Gly Gln Glu
725 730 735

Pro Leu Ala Leu Glu Ser Leu Phe Met Leu Ile Gln Gly Lys Val
740 745 750

~~Lys Val Val Asp Trp Glu Glu Ser Thr Glu Val Thr Leu Tyr Phe~~
755 760 765

5 ~~Leu Ser Leu Leu Val Phe Arg Leu Gln Asn Leu Pro Cys Gly Met~~
770 775 780

~~Glu Lys Leu Gly Ser Asp Val Ala Thr Leu Phe Thr His Ser His~~
785 790 795

10 ~~Val Val Ser Leu Val Ser Ala Ala Ala Cys Leu Leu Gly Gln Leu~~
800 805 810

15 ~~Gly Gln Gln Gly Val Thr Phe Asp Leu Gln Pro Met Glu Trp Met~~
815 820 825

~~Ala Ala Ala Thr His Ala Leu Ser Ala Pro Ala Glu Val Arg Leu~~
830 835 840

20 ~~Thr Pro Pro Gly Ser Cys Gly Phe Tyr Asp Gly Leu Leu Ile Leu~~
845 850 855

~~Leu Leu Gln Leu Leu Thr Glu Gln Gly Lys Ala Ser Leu Ile Arg~~
860 865 870

25 ~~Asp Met Ser Ser Ser Glu Met Trp Thr Val Leu Trp His Arg Phe~~
875 880 885

~~Ser Met Val Leu Arg Leu Pro Glu Glu Ala Ser Ala Gln Glu Gly~~
890 895 900

30 ~~Glu Leu Ser Leu Ser Ser Pro Pro Ser Pro Glu Pro Asp Trp Thr~~
905 910 915

35 ~~Leu Ile Ser Pro Gln Gly Met Ala Ala Leu Leu Ser Leu Ala Met~~
920 925 930

~~Ala Thr Phe Thr Gln Glu Pro Gln Leu Cys Leu Ser Cys Leu Ser~~
935 940 945

40 ~~Gln His Gly Ser Ile Leu Met Ser Ile Leu Lys His Leu Leu Cys~~
950 955 960

45 ~~Pro Ser Phe Leu Asn Gln Leu Arg Gln Ala Pro His Gly Ser Glu~~
965 970 975

~~Phe Leu Pro Val Val Leu Ser Val Cys Gln Leu Leu Cys Phe~~
980 985 990

1 Pro Phe Ala Leu Asp Met Asp Ala Asp Leu Leu Ile Val Val Leu
2 995 1000 1005

5 Ala Asp Leu Arg Asp Ser Glu Val Ala Ala His Leu Leu Gln Val
6 1010 1015 1020

10 Cys Cys Tyr His Leu Pro Leu Met Gln Val Glu Leu Pro Ile Ser
11 1025 1030 1035

15 Leu Leu Thr Arg Leu Ala Leu Met Asp Pro Thr Ser Leu Asn Gln
16 1040 1045 1050

20 Phe Val Asn Thr Val Ser Ala Ser Pro Arg Thr Ile Val Ser Phe
21 1055 1060 1065

25 Leu Ser Val Ala Leu Leu Ser Asp Gln Pro Leu Leu Thr Ser Asp
26 1070 1075 1080

30 Leu Leu Ser Leu Leu Ala His Thr Ala Arg Val Leu Ser Pro Ser
31 1085 1090 1095

35 His Leu Ser Phe Ile Gln Glu Leu Leu Ala Gly Ser Asp Glu Ser
36 1100 1105 1110

40 Tyr Arg Pro Leu Arg Ser Leu Leu Gly His Pro Glu Asn Ser Val
41 1115 1120 1125

45 Arg Ala His Thr Tyr Arg Leu Leu Gly His Leu Leu Gln His Ser
46 1130 1135 1140

50 Met Ala Leu Arg Gly Ala Leu Gln Ser Gln Ser Gly Leu Leu Ser
51 1145 1150 1155

55 Leu Leu Leu Leu Gly Leu Gly Asp Lys Asp Pro Val Val Arg Cys
56 1160 1165 1170

60 Ser Ala Ser Phe Ala Val Gly Asn Ala Ala Tyr Gln Ala Gly Pro
61 1175 1180 1185

65 Leu Gly Pro Ala Leu Ala Ala Val Pro Ser Met Thr Gln Leu
66 1190 1195 1200

70 Leu Gly Asp Pro Gln Ala Gly Ile Arg Arg Asn Val Ala Ser Ala
71 1205 1210 1215

75 Leu Gly Asn Leu Gly Pro Glu Gly Leu Gly Glu Glu Leu Leu Gln
76 1220 1225 1230

~~Cys Glu Val Pro Gln Arg Leu Leu Glu Met Ala Cys Gly Asp Pro~~
1235 1240 1245

5 ~~Gln Pro Asn Val Lys Glu Ala Ala Leu Ile Ala Leu Arg Ser Leu~~
1250 1255 1260

~~Gln Gln Glu Pro Gly Ile His Gln Val Leu Val Ser Leu Gly Ala~~
1265 1270 1275

10 ~~Ser Glu Lys Leu Ser Leu Leu Ser Leu Gly Asn Gln Ser Leu Pro~~
1280 1285 1290

~~His Ser Ser Pro Arg Pro Ala Ser Ala Lys His Cys Arg Lys Leu~~
1295 1300 1305

15 ~~Ile His Leu Leu Arg Pro Ala His Ser Met~~
1310 1315

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<211> 48

<212> DNA

<213> Artificial sequence

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<223> Artificial sequence 1 48

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<211> 48

<212> DNA

<213> Artificial sequence

40 <220>

<223> Artificial sequence 1 48

45 <400> 12

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<211> 396

<212> PRT

<213> *Mus musculus*

55 <400> 13

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60 ~~Leu Ala Leu Ser Ala Gln Ser Cys Gly Pro Gly Arg Gly Pro Val~~
20 25 30

65 ~~Gly Arg Arg Arg Tyr Val Arg Lys Gln Leu Val Pro Leu Leu Tyr~~
35 40 45

70 ~~Lys Gln Phe Val Pro Ser Met Pro Glu Arg Thr Leu Gly Ala Ser~~
50 55 60

75 ~~Gly Pro Ala Glu Gly Arg Val Thr Arg Gly Ser Glu Arg Phe Arg~~
65 70 75

Asp Leu Val Pro Asn Tyr Asn Pro Asp Ile Ile Phe Lys Asp Glu
80 85 90

5 Glu Asn Ser Gly Ala Asp Arg Leu Met Thr Glu Arg Cys Lys Glu
95 100 105

Arg Val Asn Ala Leu Ala Ile Ala Val Met Asn Met Trp Pro Gly
110 115 120

10 Val Arg Leu Arg Val Thr Glu Gly Trp Asp Glu Asp Gly His His
125 130 135

15 Ala Gln Asp Ser Leu His Tyr Glu Gly Arg Ala Leu Asp Ile Thr
140 145 150

Thr Ser Asp Arg Asp Arg Asn Lys Tyr Gly Leu Leu Ala Arg Leu
155 160 165

20 Ala Val Glu Ala Gly Phe Asp Trp Val Tyr Tyr Glu Ser Arg Asn
170 175 180

His Ile His Val Ser Val Lys Ala Asp Asn Ser Leu Ala Val Arg
185 190 195

25 Ala Gly Gly Cys Phe Pro Gly Asn Ala Thr Val Arg Leu Arg Ser
200 205 210

Gly Glu Arg Lys Gly Leu Arg Glu Leu His Arg Gly Asp Trp Val
215 220 225

30 Leu Ala Ala Asp Ala Ala Gly Arg Val Val Pro Thr Pro Val Leu
230 235 240

35 Leu Phe Leu Asp Arg Asp Leu Cln Arg Arg Ala Ser Phe Val Ala
245 250 255

Val Glu Thr Glu Arg Pro Pro Arg Lys Leu Leu Leu Thr Pro Trp
260 265 270

5 His Leu Val Phe Ala Ala Arg Gly Pro Ala Pro Ala Pro Gly Asp
275 280 285

Phe Ala Pro Val Phe Ala Arg Arg Leu Arg Ala Gly Asp Ser Val
290 295 300

10 Leu Ala Pro Gly Gly Asp Ala Leu Gln Pro Ala Arg Val Ala Arg
305 310 315

Val Ala Arg Glu Glu Ala Val Gly Val Phe Ala Pro Leu Thr Ala
320 325 330

15 His Gly Thr Leu Leu Val Asn Asp Val Leu Ala Ser Cys Tyr Ala
335 340 345

20 Val Leu Glu Ser His Gln Trp Ala His Arg Ala Phe Ala Pro Leu
350 355 360

Arg Leu Leu His Ala Leu Gly Ala Leu Leu Pro Gly Gly Ala Val
365 370 375

25 Gln Pro Thr Gly Met His Trp Tyr Ser Arg Leu Leu Tyr Arg Leu
380 385 390

Ala Glu Glu Leu Met Gly
395 396

30 <210> 14

<211> 437

<212> PRT

<213> Mus musculus

35 <400> 14

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40 Ser Leu Leu Val Cys Pro Gly Leu Ala Cys Gly Pro Gly Arg Gly
20 25 30

Phe Gly Lys Arg Arg His Pro Lys Lys Leu Thr Pro Leu Ala Tyr
35 40 45

45 Lys Gln Phe Ile Pro Asn Val Ala Glu Lys Thr Leu Gly Ala Ser
50 55 60

~~Gly Arg Tyr Glu Glu Lys Ile Thr Arg Asn Ser Glu Arg Phe Lys~~
65 70 75

5 ~~Glu Leu Thr Pro Asn Tyr Asn Pro Asp Ile Ile Phe Lys Asp Glu~~
80 85 90

~~Glu Asn Thr Gly Ala Asp Arg Leu Met Thr Gln Arg Cys Lys Asp~~
95 100 105

10 ~~Lys Leu Asn Ala Leu Ala Ile Ser Val Met Asn Gln Trp Pro Gly~~
110 115 120

15 ~~Val Arg Leu Arg Val Thr Glu Gly Trp Asp Glu Asp Gly His His~~
125 130 135

~~Ser Glu Glu Ser Leu His Tyr Glu Gly Arg Ala Val Asp Ile Thr~~
140 145 150

20 ~~Thr Ser Asp Arg Asp Arg Ser Lys Tyr Gly Met Leu Ala Arg Leu~~
155 160 165

~~Ala Val Glu Ala Gly Phe Asp Trp Val Tyr Tyr Glu Ser Lys Ala~~
170 175 180

25 ~~His Ile His Cys Ser Val Lys Ala Glu Asn Ser Val Ala Ala Lys~~
185 190 195

30 ~~Ser Gly Gly Cys Phe Pro Gly Ser Ala Thr Val His Leu Glu Gln~~
200 205 210

~~Gly Gly Thr Lys Leu Val Lys Asp Leu Arg Pro Gly Asp Arg Val~~
215 220 225

35 ~~Leu Ala Ala Asp Asp Gln Gly Arg Leu Leu Tyr Ser Asp Phe Leu~~
230 235 240

~~Thr Phe Leu Asp Arg Asp Glu Gly Ala Lys Lys Val Phe Tyr Val~~
245 250 255

40 ~~Ile Glu Thr Leu Glu Pro Arg Glu Arg Leu Leu Leu Thr Ala Ala~~
260 265 270

45 ~~His Leu Leu Phe Val Ala Pro His Asn Asp Ser Gly Pro Thr Pro~~
275 280 285

~~Gly Pro Ser Ala Leu Phe Ala Ser Arg Val Arg Pro Gly Gln Arg~~
290 295 300

Val Tyr Val Val Ala Glu Arg Gly Gly Asp Arg Arg Leu Leu Pro
305 310 315

5 Ala Ala Val His Ser Val Thr Leu Arg Glu Glu Glu Ala Gly Ala
320 325 330

Tyr Ala Pro Leu Thr Ala His Gly Thr Ile Leu Ile Asn Arg Val
335 340 345

10 Leu Ala Ser Cys Tyr Ala Val Ile Glu Glu His Ser Trp Ala His
350 355 360

15 Arg Ala Phe Ala Pro Phe Arg Leu Ala His Ala Leu Leu Ala Ala
365 370 375

Leu Ala Pro Ala Arg Thr Asp Gly Gly Gly Gly Ser Ile Pro
380 385 390

20 Ala Ala Gln Ser Ala Thr Glu Ala Arg Gly Ala Glu Pro Thr Ala
395 400 405

Gly Ile His Trp Tyr Ser Gln Leu Leu Tyr His Ile Gly Thr Trp
410 415 420

25 Leu Leu Asp Ser Glu Thr Met His Pro Leu Gly Met Ala Val Lys
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Ala Ser
437

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<212> PRT

<213> Artificial sequence

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<223> Artificial sequence 1 803

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20 25 30

Arg Gly Ala Ala Leu Ser Gly Asn Val Thr Gly Pro Gly Pro Arg
35 40 45

~~Ser Ala Gly Gly Ser Ala Arg Arg Asn Ala Pro Val Thr Ser Pro~~
50 55 60

5 ~~Pro Pro Pro Leu Leu Ser His Cys Gly Arg Ala Ala His Cys Glu~~
65 70 75

~~Pro Leu Arg Tyr Asn Val Cys Leu Gly Ser Ala Leu Pro Tyr Gly~~
80 85 90

10 ~~Ala Thr Thr Leu Leu Ala Gly Asp Ser Asp Ser Gln Glu Glu~~
95 100 105

15 ~~Ala His Ser Lys Leu Val Leu Trp Ser Gly Leu Arg Asn Ala Pro~~
110 115 120

~~Arg Cys Trp Ala Val Ile Gln Pro Leu Leu Cys Ala Val Tyr Met~~
125 130 135

20 ~~Pro Lys Cys Glu Asn Asp Arg Val Glu Leu Pro Ser Arg Thr Leu~~
140 145 150

~~Cys Gln Ala Thr Arg Gly Pro Cys Ala Ile Val Glu Arg Glu Arg~~
155 160 165

25 ~~Gly Trp Pro Asp Phe Leu Arg Cys Thr Pro Asp His Phe Pro Glu~~
170 175 180

30 ~~Gly Cys Pro Asn Glu Val Gln Asn Ile Lys Phe Asn Ser Ser Gly~~
185 190 195

~~Gln Cys Glu Ala Pro Leu Val Arg Thr Asp Asn Pro Lys Ser Trp~~
200 205 210

35 ~~Tyr Glu Asp Val Glu Gly Cys Gly Ile Cln Cys Gln Asn Pro Leu~~
215 220 225

~~Phe Thr Glu Ala Glu His Gln Asp Met His Ser Tyr Ile Ala Ala~~
230 235 240

40 ~~Phe Gly Ala Val Thr Gly Leu Cys Thr Leu Phe Thr Leu Ala Thr~~
245 250 255

45 ~~Phe Val Ala Asp Trp Arg Asn Ser Asn Arg Tyr Pro Ala Val Ile~~
260 265 270

~~Leu Phe Tyr Val Asn Ala Cys Phe Phe Val Gly Ser Ile Gly Trp~~
275 280 285

Leu Ala Cln Phe Met Asp Gly Ala Arg Arg Glu Ile Val Cys Arg
290 295 300

5 Ala Asp Gly Thr Met Arg Phe Gly Glu Pro Thr Ser Ser Glu Thr
305 310 315

Leu Ser Cys Val Ile Ile Phe Val Ile Val Tyr Tyr Ala Leu Met
320 325 330

10 Ala Gly Val Val Trp Phe Val Val Leu Thr Tyr Ala Trp His Thr
335 340 345

Ser Phe Lys Ala Leu Gly Thr Thr Tyr Cln Pro Leu Ser Gly Lys
350 355 360

15 Thr Ser Tyr Phe His Leu Leu Thr Trp Ser Leu Pro Phe Val Leu
365 370 375

Thr Val Ala Ile Leu Ala Val Ala Cln Val Asp Gly Asp Ser Val
20 380 385 390

Ser Gly Ile Cys Phe Val Gly Tyr Lys Asn Tyr Arg Tyr Arg Ala
395 400 405

25 Gly Phe Val Leu Ala Pro Ile Gly Leu Val Leu Ile Val Gly Gly
410 415 420

Tyr Phe Leu Ile Arg Gly Val Met Thr Leu Phe Ser Ile Lys Ser
425 430 435

30 Asn His Pro Gly Leu Leu Ser Glu Lys Ala Ala Ser Lys Ile Asn
440 445 450

Glu Thr Met Leu Arg Leu Gly Ile Phe Gly Phe Leu Ala Phe Gly
35 455 460 465

Phe Val Leu Ile Thr Phe Ser Cys His Phe Tyr Asp Phe Phe Asn
470 475 480

40 Cln Ala Glu Trp Glu Arg Ser Phe Arg Asp Tyr Val Leu Cys Cln
485 490 495

Ala Asn Val Thr Ile Gly Leu Pro Thr Lys Lys Pro Ile Pro Asp
45 500 505 510

Cys Glu Ile Lys Asn Arg Pro Ser Leu Leu Val Glu Lys Ile Asn
515 520 525

Leu Phe Ala Met Phe Gly Thr Gly Ile Ala Met Ser Thr Trp Val
530 535 540

5 Trp Thr Lys Ala Thr Leu Leu Ile Trp Arg Arg Thr Trp Cys Arg
545 550 555

Leu Thr Gly His Ser Asp Asp Glu Pro Lys Arg Ile Lys Lys Ser
10 560 565 570

15 Lys Met Ile Ala Lys Ala Phe Ser Lys Arg Arg Glu Leu Leu Cln
575 580 585

Asn Pro Gly Cln Glu Leu Ser Phe Ser Met His Thr Val Ser His
15 590 595 600

20 Asp Gly Pro Val Ala Gly Leu Ala Phe Glu Leu Asn Glu Pro Ser
605 610 615

Ala Asp Val Ser Ser Ala Trp Ala Gln His Val Thr Lys Met Val
25 620 625 630

Ala Arg Arg Gly Ala Ile Leu Pro Gln Asp Val Ser Val Thr Pro
30 635 640 645

Val Ala Thr Pro Val Pro Pro Glu Glu Gln Ala Asn Leu Trp Leu
35 650 655 660

Val Glu Ala Glu Ile Ser Pro Glu Leu Glu Lys Arg Leu Gly Arg
40 665 670 675

Lys Lys Lys Arg Arg Lys Arg Lys Lys Glu Val Cys Pro Leu Gly
45 680 685 690

Pro Ala Pro Glu Leu His His Ser Ala Pro Val Pro Ala Thr Ser
50 695 700 705

Ala Val Pro Arg Leu Pro Gln Leu Pro Arg Gln Lys Cys Leu Val
55 710 715 720

Ala Ala Asn Ala Trp Gly Thr Gly Glu Pro Cys Arg Cln Gly Ala
60 725 730 735

Trp Thr Val Val Ser Asn Pro Phe Cys Pro Glu Pro Ser Pro His
65 740 745 750

70 Gln Asp Pro Phe Leu Pro Gly Ala Ser Ala Pro Arg Val Trp Ala
755 760 765

~~Cln Gly Arg Leu Cln Gly Leu Gly Ser Ile His Ser Arg Thr Asn~~
770 775 780

5 ~~Leu Met Glu Ala Glu Leu Leu Asp Ala Asp Ser Asp Phe Glu Cln~~
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~~Lys Leu Ile Ser Glu Glu Asp Leu~~
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25 ~~Arg Gly Ala Ala Leu Ser Gly Asn Val Thr Gly Pro Gly Pro Arg~~
35 40 45

~~Ser Ala Gly Gly Ser Ala Arg Arg Asn Ala Pro Val Thr Ser Pro~~
50 55 60

30 ~~Pro Pro Pro Leu Leu Ser His Cys Gly Arg Ala Ala His Cys Glu~~
65 70 75

35 ~~Pro Leu Arg Tyr Asn Val Cys Leu Gly Ser Ala Leu Pro Tyr Gly~~
80 85 90

~~Ala Thr Thr Thr Leu Leu Ala Gly Asp Ser Asp Ser Gln Glu Glu~~
95 100 105

40 ~~Ala His Ser Lys Leu Val Leu Trp Ser Gly Leu Arg Asn Ala Pro~~
110 115 120

~~Arg Cys Trp Ala Val Ile Gln Pro Leu Leu Cys Ala Val Tyr Met~~
125 130 135

45 ~~Pro Lys Cys Glu Asn Asp Arg Val Glu Leu Pro Ser Arg Thr Leu~~
140 145 150

~~Cys Gln Ala Thr Arg Gly Pro Cys Ala Ile Val Glu Arg Glu Arg~~
155 160 165

5 ~~Gly Trp Pro Asp Phe Leu Arg Cys Thr Pro Asp His Phe Pro Glu~~
170 175 180

~~Gly Cys Pro Asn Glu Val Gln Asn Ile Lys Phe Asn Ser Ser Gly~~
185 190 195

10 ~~Gln Cys Glu Ala Pro Leu Val Arg Thr Asp Asn Pro Lys Ser Trp~~
200 205 210

15 ~~Tyr Glu Asp Val Glu Gly Cys Gly Ile Gln Cys Gln Asn Pro Leu~~
215 220 225

~~Phe Thr Glu Ala Glu His Gln Asp Met His Ser Tyr Ile Ala Ala~~
230 235 240

20 ~~Phe Gly Ala Val Thr Gly Leu Cys Thr Leu Phe Thr Leu Ala Thr~~
245 250 255

~~Phe Val Ala Asp Trp Arg Asn Ser Asn Arg Tyr Pro Ala Val Ile~~
260 265 270

25 ~~Leu Phe Tyr Val Asn Ala Cys Phe Phe Val Gly Ser Ile Gly Trp~~
275 280 285

~~Leu Ala Gln Phe Met Asp Gly Ala Arg Arg Glu Ile Val Cys Arg~~
290 295 300

30 ~~Ala Asp Gly Thr Met Arg Phe Gly Glu Pro Thr Ser Ser Glu Thr~~
305 310 315

35 ~~Leu Ser Cys Val Ile Ile Phe Val Ile Val Tyr Tyr Ala Leu Met~~
320 325 330

~~Ala Gly Val Val Trp Phe Val Val Leu Thr Tyr Ala Trp His Thr~~
335 340 345

40 ~~Ser Phe Lys Ala Leu Gly Thr Thr Tyr Gln Pro Leu Ser Gly Lys~~
350 355 360

45 ~~Thr Ser Tyr Phe His Leu Leu Thr Trp Ser Leu Pro Phe Val Leu~~
365 370 375

~~Thr Val Ala Ile Leu Ala Val Ala Gln Val Asp Gly Asp Ser Val~~
380 385 390

~~Ser Gly Ile Cys Phe Val Gly Tyr Lys Asn Tyr Arg Tyr Arg Ala~~
395 400 405

5 ~~Gly Phe Val Leu Ala Pro Ile Gly Leu Val Leu Ile Val Gly Gly~~
410 415 420

~~Tyr Phe Leu Ile Arg Gly Val Met Thr Leu Phe Ser Ile Lys Ser~~
425 430 435

10 ~~Asn His Pro Gly Leu Leu Ser Glu Lys Ala Ala Ser Lys Ile Asn~~
440 445 450

15 ~~Glu Thr Met Leu Arg Leu Gly Ile Phe Gly Phe Leu Ala Phe Gly~~
455 460 465

~~Phe Val Leu Ile Thr Phe Ser Cys His Phe Tyr Asp Phe Phe Asn~~
470 475 480

20 ~~Gln Ala Glu Trp Glu Arg Ser Phe Arg Asp Tyr Val Leu Cys Gln~~
485 490 495

~~Ala Asn Val Thr Ile Gly Leu Pro Thr Lys Lys Pro Ile Pro Asp~~
500 505 510

25 ~~Cys Glu Ile Lys Asn Arg Pro Ser Leu Leu Val Glu Lys Ile Asn~~
515 520 525

30 ~~Leu Phe Ala Met Phe Gly Thr Gly Ile Ala Met Ser Thr Leu Val~~
530 535 540

~~Trp Thr Lys Ala Thr Leu Leu Ile Trp Arg Arg Thr Trp Cys Arg~~
545 550 555

35 ~~Leu Thr Gly His Ser Asp Asp Glu Pro Lys Arg Ile Lys Lys Ser~~
560 565 570

~~Lys Met Ile Ala Lys Ala Phe Ser Lys Arg Arg Glu Leu Leu Gln~~
575 580 585

40 ~~Asn Pro Gly Gln Glu Leu Ser Phe Ser Met His Thr Val Ser His~~
590 595 600

45 ~~Asp Gly Pro Val Ala Gly Leu Ala Phe Glu Leu Asn Glu Pro Ser~~
605 610 615

~~Ala Asp Val Ser Ser Ala Trp Ala Gln His Val Thr Lys Met Val~~
620 625 630

Ala Arg Arg Gly Ala Ile Leu Pro Gln Asp Val Ser Val Thr Pro
635 640 645

5 Val Ala Thr Pro Val Pro Pro Glu Glu Gln Ala Asn Leu Trp Leu
650 655 660

Val Glu Ala Glu Ile Ser Pro Glu Leu Glu Lys Arg Leu Gly Arg
665 670 675

10 Lys Lys Lys Arg Arg Lys Arg Lys Lys Glu Val Cys Pro Leu Gly
680 685 690

15 Pro Ala Pro Glu Leu His His Ser Ala Pro Val Pro Ala Thr Ser
695 700 705

Ala Val Pro Arg Leu Pro Gln Leu Pro Arg Gln Lys Cys Leu Val
710 715 720

20 Ala Ala Asn Ala Trp Gly Thr Gly Glu Pro Cys Arg Gln Gly Ala
725 730 735

Trp Thr Val Val Ser Asn Pro Phe Cys Pro Glu Pro Ser Pro His
740 745 750

25 Gln Asp Pro Phe Leu Pro Gly Ala Ser Ala Pro Arg Val Trp Ala
755 760 765

Cln Gly Arg Leu Cln Gly Leu Gly Ser Ile His Ser Arg Thr Asn
770 775 780

30 Leu Met Glu Ala Glu Leu Leu Asp Ala Asp Ser Asp Phe
785 790 793

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Arg Leu Leu Gln Leu Leu Leu Val Leu Leu Gly Gly Arg Gly
20 25 30

45 Arg Gly Ala Ala Leu Ser Gly Asn Val Thr Gly Pro Gly Pro Arg
35 40 45

Substitute Specification

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~~Ser Ala Gly Gly Ser Ala Arg Arg Asn Ala Pro Val Thr Ser Pro~~
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5 ~~Pro Pro Pro Leu Leu Ser His Cys Gly Arg Ala Ala His Cys Glu~~
65 70 75

10 ~~Pro Leu Arg Tyr Asn Val Cys Leu Gly Ser Ala Leu Pro Tyr Gly~~
80 85 90

15 ~~Ala Thr Thr Thr Leu Leu Ala Gly Asp Ser Asp Ser Gln Glu Glu~~
95 100 105

20 ~~Ala His Ser Lys Leu Val Leu Trp Ser Gly Leu Arg Asn Ala Pro~~
110 115 120

25 ~~Arg Cys Trp Ala Val Ile Gln Pro Leu Leu Cys Ala Val Tyr Met~~
125 130 135

30 ~~Pro Lys Cys Glu Asn Asp Arg Val Glu Leu Pro Ser Arg Thr Leu~~
140 145 150

35 ~~Cys Gln Ala Thr Arg Gly Pro Cys Ala Ile Val Glu Arg Glu Arg~~
155 160 165

40 ~~Gly Trp Pro Asp Phe Leu Arg Cys Thr Pro Asp His Phe Pro Glu~~
170 175 180

45 ~~Gly Cys Pro Asn Glu Val Gln Asn Ile Lys Phe Asn Ser Ser Gly~~
185 190 195

50 ~~Gln Cys Glu Ala Pro Leu Val Arg Thr Asp Asn Pro Lys Ser Trp~~
200 205 210

55 ~~Tyr Glu Asp Val Glu Gly Cys Gly Ile Gln Cys Gln Asn Pro Leu~~
215 220 225

60 ~~Phe Thr Glu Ala Glu His Gln Asp Met His Ser Tyr Ile Ala Ala~~
230 235 240

65 ~~Phe Gly Ala Val Thr Gly Leu Cys Thr Leu Phe Thr Leu Ala Thr~~
245 250 255

70 ~~Phe Val Ala Asp Trp Arg Asn Ser Asn Arg Tyr Pro Ala Val Ile~~
260 265 270

75 ~~Leu Phe Tyr Val Asn Ala Cys Phe Phe Val Gly Ser Ile Gly Trp~~
275 280 285

Leu Ala Gln Phe Met Asp Gly Ala Arg Arg Glu Ile Val Cys Arg
290 295 300

5 Ala Asp Gly Thr Met Arg Phe Gly Glu Pro Thr Ser Ser Glu Thr
305 310 315

Leu Ser Cys Val Ile Ile Phe Val Ile Val Tyr Tyr Ala Leu Met
220 325 330

10 Ala Gly Val Val Trp Phe Val Val Leu Thr Tyr Ala Trp His Thr
335 340 345

Ser Phe Lys Ala Leu Gly Thr Thr Tyr Cln Pro Leu Ser Gly Lys
350 355 360

15 Thr Ser Tyr Phe His Leu Leu Thr Trp Ser Leu Pro Phe Val Leu
365 370 375

20 Thr Val Ala Ile Leu Ala Val Ala Gln Val Asp Gly Asp Ser Val
380 385 390

Ser Gly Ile Cys Phe Val Gly Tyr Lys Asn Tyr Arg Tyr Arg Ala
395 400 405

25 Gly Phe Val Leu Ala Pro Ile Gly Leu Val Leu Ile Val Gly Gly
410 415 420

Tyr Phe Leu Ile Arg Gly Val Met Thr Leu Phe Ser Ile Lys Ser
425 430 435

30 Asn His Pro Gly Leu Leu Ser Glu Lys Ala Ala Ser Lys Ile Asn
440 445 450

35 Glu Thr Met Leu Arg Leu Gly Ile Phe Gly Phe Leu Ala Phe Gly
455 460 465

Phe Val Leu Ile Thr Phe Ser Cys His Phe Tyr Asp Phe Phe Asn
470 475 480

40 Gln Ala Glu Trp Glu Arg Ser Phe Arg Asp Tyr Val Leu Cys Gln
485 490 495

45 Ala Asn Val Thr Ile Gly Leu Pro Thr Lys Lys Pro Ile Pro Asp
500 505 510

Cys Glu Ile Lys Asn Arg Pro Ser Leu Leu Val Glu Lys Ile Asn
515 520 525

Leu Phe Ala Met Phe Gly Thr Gly Ile Ala Met Ser Thr Trp Val
530 535 540

5 Trp Thr Lys Ala Thr Leu Leu Ile Trp Arg Arg Thr Trp Cys Arg
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Leu Thr Gly His Ser Asp Asp Glu Pro Lys Arg Ile Lys Lys Ser
10 560 565 570

15 Lys Met Ile Ala Lys Ala Phe Ser Lys Arg Arg Glu Leu Leu Gln
575 580 585

Asn Pro Gly Gln Glu Leu Ser Phe Ser Met His Thr Val Ser His
15 590 595 600

20 Asp Gly Pro Val Ala Gly Leu Ala Phe Glu Leu Asn Glu Pro Ser
605 610 615

Ala Asp Val Ser Ser Ala Trp Ala Gln His Val Thr Lys Met Val
25 620 625 630

Ala Arg Arg Gly Ala Ile Leu Pro Gln Asp Val Ser Val Thr Pro
30 635 640 645

Val Ala Thr Pro Val Pro Pro Glu Glu Gln Ala Asn Leu Trp Leu
35 650 655 660

Val Glu Ala Glu Ile Ser Pro Glu Leu Glu Lys Arg Leu Gly Arg
40 665 670 675

Lys Lys Lys Arg Arg Lys Arg Lys Lys Glu Val Cys Pro Leu Gly
45 680 685 690

Pro Ala Pro Glu Leu His His Ser Ala Pro Val Pro Ala Thr Ser
50 695 700 705

Ala Val Pro Arg Leu Pro Gln Leu Pro Arg Gln Lys Cys Leu Val
55 710 715 720

Ala Ala Asn Ala Trp Gly Thr Gly Glu Pro Cys Arg Gln Gly Ala
60 725 730 735

Trp Thr Val Val Ser Asn Pro Phe Cys Pro Glu Pro Ser Pro His
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Gln Asp Pro Phe Leu Pro Gly Ala Ser Ala Pro Arg Val Trp Ala
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15 Ser Gln Pro Arg Ala His Leu Ser Ala His Gln Ala Pro Ser Pro

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Ala Ala Leu Pro Gly Tyr Pro Ala Met Ser Pro Ala Trp Leu Arg
35 40 45

20 Pro Arg Leu Arg Phe Cys Leu Phe Leu Leu Leu Leu Leu Val

50 55 60

Pro Ala Ala Arg Gly Cys Gly Pro Gly Arg Val Val Gly Ser Arg
65 70 75

25 Arg Arg Pro Pro Arg Lys Leu Val Pro Leu Ala Tyr Lys Gln Phe

80 85 90

30 Ser Pro Asn Val Pro Glu Lys Thr Leu Gly Ala Ser Gly Arg Tyr

95 100 105

Glu Gly Lys Ile Ala Arg Ser Ser Glu Arg Phe Lys Glu Leu Thr
110 115 120

35 Pro Asn Tyr Asn Pro Asp Ile Ile Phe Lys Asp Glu Glu Asn Thr

125 130 135

Gly Ala Asp Arg Leu Met Thr Gln Arg Cys Lys Asp Arg Leu Asn
140 145 150

40 Ser Leu Ala Ile Ser Val Met Asn Gln Trp Pro Gly Val Lys Leu

155 160 165

Arg Val Thr Glu Gly Trp Asp Glu Asp Gly His His Ser Glu Glu
170 175 180

45 Ser Leu His Tyr Glu Gly Arg Ala Val Asp Ile Thr Thr Ser Asp

185 190 195

50 Arg Asp Arg Asn Lys Tyr Gly Leu Leu Ala Arg Leu Ala Val Glu

200 205 210

Ala Gly Phe Asp Trp Val Tyr Tyr Glu Ser Lys Ala His Val His
215 220 225

55 Cys Ser Val Lys Ser Glu His Ser Ala Ala Ala Lys Thr Gly Gly

230 235 240

60 Cys Phe Pro Ala Gly Ala Cln Val Arg Leu Glu Asn Gly Glu Arg

245 250 255

Val Ala Leu Ser Ala Val Lys Pro Gly Asp Arg Val Leu Ala Met
260 265 270

5 Gly Glu Asp Gly Thr Pro Thr Phe Ser Asp Val Leu Ile Phe Leu
275 280 285

Asp Arg Glu Pro Asn Arg Leu Arg Ala Phe Gln Val Ile Glu Thr
290 295 300

10 Gln Asp Pro Pro Arg Leu Ala Leu Thr Pro Ala His Leu Leu
305 310 315

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